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(54) **METHODS OF MAKING  
PHARMACEUTICAL COMPOSITIONS WITH  
MINICELLS**

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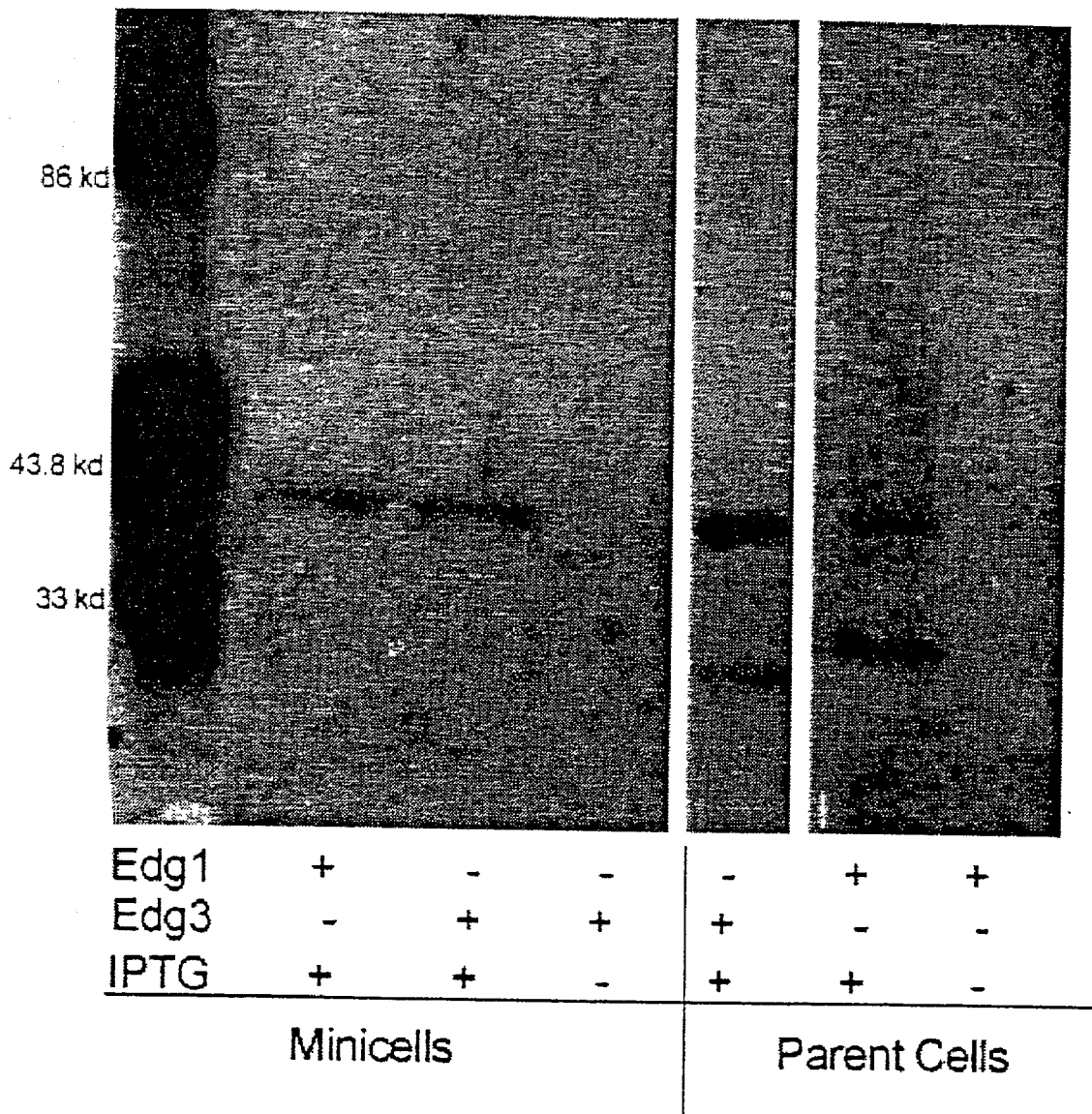
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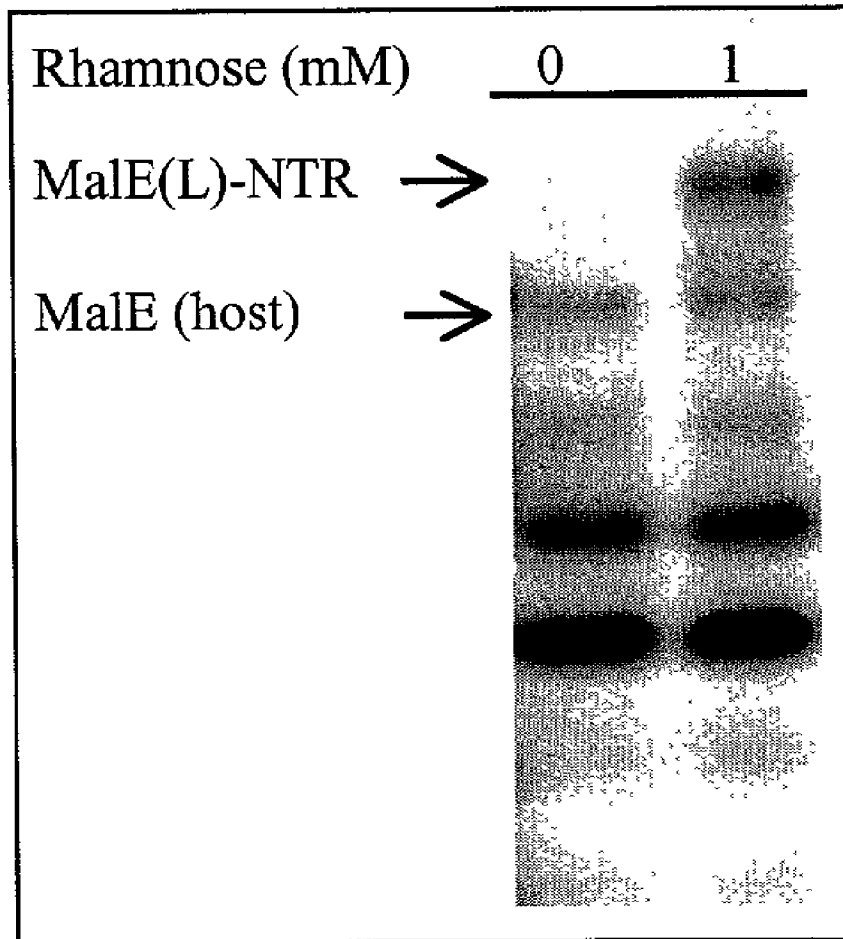
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(57) **ABSTRACT**

The invention provides compositions and methods for the production of achromosomal and anucleate cells useful for applications such as diagnostic and therapeutic uses, as well as research tools and agents for drug discovery.



**Figure 1**



**Figure 2**

## METHODS OF MAKING PHARMACEUTICAL COMPOSITIONS WITH MINICELLS

### RELATED APPLICATIONS

[0001] This application claims priority to the following U.S. patent applications:

[0002] Serial No. 60/295,566 entitled "Minicell Compositions and Methods" by Roger Sabbadini, filed May 24, 2001;

[0003] Serial No. 60/359,843 entitled "Minicell Compositions and Methods" by Sabbadini, et al., filed Feb. 25, 2002; and

[0004] Ser. No. \_\_\_\_\_ (attorney docket No. 089608-0401), entitled "Methods of Making Minicells," by Surber, et al., filed May 24, 2002.

[0005] All of the preceding applications are hereby incorporated in their entirety (including drawings) by reference thereto.

### FIELD OF THE INVENTION

[0006] The invention is drawn to compositions and methods for the production of achromosomal archeobacterial, eubacterial and anucleate eukaryotic cells that are used as, e.g., therapeutics and/or diagnostics, reagents in drug discovery and functional proteomics, research tools, and in other applications as well.

### BACKGROUND OF THE INVENTION

[0007] The following description of the background of the invention is provided to aid in understanding the invention, but is not admitted to describe or constitute prior art to the invention. The contents of the articles, patents, and patent applications, and all other documents and electronically available information mentioned or cited in this application, are hereby incorporated by reference in their entirety to the same extent as if each individual publication was specifically and individually indicated to be incorporated by reference. Applicants reserve the right to physically incorporate into this application any and all materials and information from any such articles, patents, patent applications, or other documents.

[0008] Minicells are achromosomal cells that are products of aberrant cell division, and contain RNA and protein, but little or no chromosomal DNA. Clark-Curtiss and Curtiss III, Analysis of Recombinant DNA Using *Escherichia coli* Minicells, 101 Methods in Enzymology 347 (1983); Reeve and Mendelson, Minicells of *Bacillus subtilis*. A new system for transport studies in absence of macromolecular biosynthesis, 352 Biochim. Biophys. Acta 298-305 (1974). Minicells are capable of plasmid-directed synthesis of discrete polypeptides in the absence of synthesis directed by mRNA from the bacterial chromosome. Meagher et al., Protein Expression in *E. coli* Minicells by Recombinant Plasmids, 10 Cell 521, 523 (1977); Roozen et al., Synthesis of Ribonucleic Acid and Protein in Plasmid-Containing Minicells of *Escherichia coli* K-12, 107(1) J. of Bacteriology 21 (1971); and Curtiss III, Research on bacterial conjugation with minicells and minicell-producing *E. coli* strains, In: Microbial Drug Resistance, Editors Susumu Mitsuhashi & Hajime Hashimoto, p. 169 (Baltimore: University Park Press 1976).

Early descriptions of minicells include those of Adler et al., Genetic control of cell division in bacteria, 154 Science 417 (1966), and Adler et al. (Miniature *Escherichia coli* cells deficient in DNA, 57 Proc. Nat. Acad. Sci (Wash.) 321 (1967)). However, discovery of the production of minicells can arguably be traced to the 1930's (Frazer and Curtiss III, Production, Properties and Utility of Bacterial Minicells, 69 Curr. Top. Microbiol. Immunol. 1-3 (1975)).

[0009] Prokaryotic (a.k.a. eubacterial) minicells have been used to produce various eubacterial proteins. See, e.g., Michael Gaael, et al., The kdpF Subunit Is Part of the K<sup>+</sup>-translocating Kdp Complex of *Escherichia coli* and Is Responsible for Stabilization of the Complex in vitro, 274(53) Jn. of Biological Chemistry 37901 (1999); Harlow, et al., Cloning and Characterization of the gsk Gene Encoding Guanosine Kinase of *Escherichia coli*, 177(8) J. of Bacteriology 2236 (1995); Carol L. Pickett, et al., Cloning, Sequencing, and Expression of the *Escherichia coli* Cytolethal Distinding Toxin Genes, 62(3) Infection & Immunity 1046 (1994); Raimund Eck & Jorn Belter, Cloning and characterization of a gene coding for the catechol 1,2 dioxygenase of *Arthrobacter* sp. mA3, 123 Gene 87 (1993); Andreas Schlösser, et al, Subcloning, Nucleotide Sequence, and Expression of trkG, a Gene That Encodes an Integral Membrane Protein Involved in Potassium Uptake via the Trk System of *Escherichia coli*, 173(10) J. of Bacteriology 3170 (1991); Mehrdad Jannatipour, et al., Translocation of *Vibrio harveyi* N,N'-Diacetylchitinase to the Outer Membrane of *Escherichia coli* 169(8) J. of Bacteriology 3785 (1987); and Jacobs et al., Expression of *Mycobacterium leprae* genes from a *Streptococcus mutans* promoter in *Escherichia coli* K-12, 83(6) Proc. Natl. Acad. Sci. USA 1926 (1986);

[0010] Various bacteria have been used, or proposed to be used, as gene delivery vectors to mammalian cells. For reviews, see Grillot-Courvalin et al., Bacteria as gene delivery vectors for mammalian cells, 10 Current Opinion in Biotechnology 477 (1999); Johnsen et al., Transfer of DNA from Genetically Modified Organisms (GMOs), Biotechnological Institute, 1-70 (2000); Sizemore et al., Attenuated *Shigella* as a DNA delivery vehicle for DNA-mediated immunization, 270(5234) Science 299 (1995); Patrice Courvalin, et al., Gene transfer from bacteria to mammalian cells, 318 C. R. Acad. Sci. 1207 (1995); Sizemore, et al. Attenuated bacteria as a DNA delivery vehicle for DNA-mediated immunization, 15(8) Vaccine 804 (1997).

[0011] U.S. Pat. No. 4,190,495, which issued Feb. 26, 1980, to Curtiss is drawn to minicell producing strains of *E. coli* that are stated to be useful for the recombinant expression of proteins.

[0012] U.S. Pat. No. 4,311,797, which issued Jan. 19, 1982 to Khachatourians is stated to be drawn to a minicell based vaccine. The vaccine is stated to induce the production of antibodies against enteropathogenic *E. coli* cells in cattle and is stated to be effective against coliform enteritis.

[0013] Eubacterial minicells expressing immunogens from other prokaryotes have been described. Purcell et al., Molecular cloning and characterization of the 15-kilodalton major immunogen of *Treponema pallidum*, Infect. Immun. 57:3708, 1989.

[0014] In "Biotechnology: Promise . . . and Peril" (IDRC Reports 9:4-7, 1980) authors Fleury and Shirkie aver that



George Khachatourians at the University of Saskatchewan, Canada, "is working on a vaccine against cholera using 'minicells.'" The minicells are said to contain "genes from the pathogenic agent," and the "pathogen antigens are carried on the surface of the minicells" (p. 5, paragraph bridging the central and right columns).

[0015] Lundstrom et al., Secretion of Semliki Forest virus membrane glycoprotein E1 from *Bacillus subtilis*, *Virus Res.* 2:69-83, 1985, describe the expression of the E1 protein of the eukaryotic virus, Semliki Forest virus (SFV), in *Bacillus* minicells. The SFV E1 protein used in these studies is not the native E1 protein. Rather, it is a fusion protein in which the N-terminal signal sequence and C-terminal transmembrane domain have been removed and replaced with signal sequences from a gene from *Bacillus amyloliquefaciens*. The authors aver that "E1 is properly translocated through the cell membrane and secreted" (p. 81, 1.1.19-20), and note that "it has been difficult to express viral membrane proteins in prokaryotes" (p. 81, 1.27).

[0016] U.S. Pat. No. 4,237,224, which issued Dec. 2, 1980, to Cohen and Boyer, describes the expression of *X. Laevis* DNA in *E. coli* minicells.

[0017] U.S. patent application Serial No. 60/293,566 (attorney docket Nos. 078853-0401 and 089608-0201), is entitled "Minicell Compositions and Methods," and was filed May 24, 2001, by Sabbadini, Roger A., Berkley, Neil L., and Klepper, Robert E., and is hereby incorporated in its entirety by reference.

[0018] Jespersen et al. describes the use of "proteoliposomes" to generate antibodies to the AMPA receptor. Jespersen L K, Kuusinen A, Orellana A, Keinanen K, Engberg J. Use of proteoliposomes to generate phage antibodies against native AMPA receptor. *Eur J Biochem.* 2000 March;267(5): 1382-9

#### SUMMARY OF THE INVENTION

[0019] The invention is drawn to compositions and methods for the production and use of minicells, including but not limited to eubacterial minicells, in applications such as diagnostics, therapeutics, research, compound screening and drug discovery, as well as agents for the delivery of nucleic acids and other bioactive compounds to cells.

[0020] Minicells are derivatives of cells that lack chromosomal DNA and which are sometimes referred to as anucleate cells. Because eubacterial and archeobacterial cells, unlike eukaryotic cells, do not have a nucleus (a distinct organelle that contains chromosomes), these non-eukaryotic minicells are more accurately described as being "without chromosomes" or "achromosomal," as opposed to "anucleate." Nonetheless, those skilled in the art often use the term "anucleate" when referring to bacterial minicells in addition to other minicells. Accordingly, in the present disclosure, the term "minicells" encompasses derivatives of eubacterial cells that lack a chromosome; derivatives of archeobacterial cells that lack their chromosome(s), and anucleate derivatives of eukaryotic cells. It is understood, however, that some of the relevant art may use the terms "anucleate minicells" or "anucleate cells" loosely to refer to any of the preceding types of minicells.

[0021] In one aspect, the invention is drawn to a eubacterial minicell comprising a membrane protein that is not

naturally found in a prokaryote, i.e., a membrane protein from a eukaryote or an archaeobacterium. Such minicells may, but need not, comprise an expression element that encodes and expresses the membrane protein that it comprises. The membrane protein may be one found in any non-eubacterial membrane, including, by way of non-limiting example, a cellular membrane, a nuclear membrane, a nucleolar membrane, a membrane of the endoplasmic reticulum (ER), a membrane of a Golgi body, a membrane of a lysosome, a membrane of a peroxisome, a caveolar membrane, an outer membrane of a mitochondrion or a chloroplast, and an inner membrane of a mitochondrion or a chloroplast. By way of non-limiting example, a membrane protein may be a receptor, such as a G-protein coupled receptor; an enzyme, such as ATPase or adenylate cyclase, a cytochrome; a channel; a transporter; or a membrane-bound nucleic acid binding factor, such as a transcription and/or translation factor; signaling components; components of the electron transport chain (ETC); or cellular antigens. A membrane fusion protein, which is generated in vitro using molecular cloning techniques, does not occur in nature and is thus a membrane protein that is not naturally found in a prokaryote, even if the fusion protein is prepared using amino acid sequences derived from eubacterial proteins.

[0022] Minicells that have segregated from parent cells lack chromosomal and/or nuclear components, but retain the cytoplasm and its contents, including the cellular machinery required for protein expression. Although chromosomes do not segregate into minicells, extrachromosomal and/or episomal genetic expression elements will segregate, or may be introduced into minicells after segregation from parent cells. Thus, in one aspect, the invention is drawn to minicells comprising an expression element, which may be an inducible expression element, that comprises expression sequences operably linked to an open reading frame (ORF) that encodes the non-eubacterial membrane protein. In a related aspect, the invention is drawn to minicell-producing host cells having an expression element, which may be an inducible expression element, that comprises expression sequences operably linked to an ORF that encodes a non-eubacterial membrane protein. In a related aspect, the invention is drawn to a method of making a eubacterial minicell comprising a membrane protein that is not naturally found in a prokaryote, the method comprising growing minicell-producing host cells, the host cells having an expression element, which may be an inducible expression element, that comprises expression sequences operably linked to an ORF that encodes a non-eubacterial membrane protein; and preparing minicells from the host cells. Optionally, at any point in the method, an inducing agent is provided in order to induce expression of an ORF that encodes a non-eubacterial membrane protein.

[0023] In one aspect, the invention is drawn to display produced membrane-associated protein(s) on the surface of the minicell. For purposes of this document, the term "display" is defined as exposure of the structure of interest on the outer surface of the minicell. By way of non-limiting example, this structure may be an internally expressed membrane protein or chimeric construct to be inserted in or associated with the minicell membrane such that the extracellular domain or domain of interest is exposed on the outer surface of the minicell (expressed and displayed on the surface of the minicell or expressed in the parental cell to be displayed on the surface of the segregated minicell). In any

scenario, the “displayed” protein or protein domain is available for interaction with extracellular components. A membrane-associated protein may have more than one extracellular domain, and a minicell of the invention may display more than one membrane-associated protein.

**[0024]** A membrane protein displayed by eubacterial minicells may be a receptor. Receptors include, by way of non-limiting example, G-coupled protein receptors, hormone receptors, and growth factor receptors. Minicells displaying a receptor may, but need not, bind ligands of the receptor. In therapeutic applications of this aspect of the invention, the ligand is an undesirable compound that is bound to its receptor and, in some aspects, is internalized or inactivated by the minicells. In drug discovery applications of this aspect of the invention, the ligand for the receptor may be detectably labeled so that its binding to its receptor may be quantified. In the latter circumstance, the minicells may be used to identify and isolate, from a pool of compounds, one or more compounds that inhibit or stimulate the activity of the receptor. That is, these minicells can be used in screening assays, including assays such as those used in high throughput screening (HTS) systems and other drug discovery methods, for the purpose of preparing compounds that influence the activity of a receptor of interest.

**[0025]** The displayed domain of a membrane protein may be an enzymatic domain such as on having oxidoreductase, transferase, hydrolase, lyase, isomerase ligase, lipase, kinase, phosphatase, protease, nuclease and/or synthetase activity. Contacting such minicells with the appropriate substrate of the enzyme allows the substrate to be converted to reactant. When either the substrate or reactant is detectable, the reaction catalyzed by the membrane-bound enzyme may be quantified. In the latter instance, the minicells may be used to identify and isolate, from a pool of compounds, one or more compounds that inhibit or stimulate the activity of the enzyme represented by the displayed enzymatic moiety. That is, these minicells can be used in screening assays, including assays such as those used in high throughput screening (HTS) systems and other drug discovery methods, for the purpose of preparing compounds that influence the activity of an enzyme or enzymatic moiety of interest.

**[0026]** The membrane protein displayed by minicells may be a fusion protein, i.e., a protein that comprises a first polypeptide having a first amino acid sequence and a second polypeptide having a second amino acid sequence, wherein the first and second amino acid sequences are not naturally present in the same polypeptide. At least one polypeptide in a membrane fusion protein is a “transmembrane domain” or “membrane-anchoring domain”. The transmembrane and membrane-anchoring domains of a membrane fusion protein may be selected from membrane proteins that naturally occur in a eucaryote, such as a fungus, a unicellular eucaryote, a plant and an animal, such as a mammal including a human. Such domains may be from a viral membrane protein naturally found in a virus such as a bacteriophage or a eucaryotic virus, e.g., an adenovirus or a retrovirus. Such domains may be from a membrane protein naturally found in an archaebacterium such as a thermophile.

**[0027]** The displayed domain of a membrane fusion protein may be an enzymatic domain such as one having oxidoreductase, transferase, hydrolase, lyase, isomerase

ligase, lipase, kinase, phosphatase, protease, nuclease and/or synthetase activity. Contacting such minicells with the appropriate substrate of the enzyme allows the substrate to be converted to reactant. When either the substrate or reactant is detectable, the reaction catalyzed by the membrane-bound enzyme may be quantified. In the latter instance, the minicells may be used to identify and isolate, from a pool of compounds, one or more compounds that inhibit or stimulate the activity of the enzyme represented by the displayed enzymatic moiety. That is, these minicells can be used in screening assays, including assays such as those used in high throughput screening (HTS) systems and other drug discovery methods, for the purpose of preparing compounds that influence the activity of an enzyme or enzymatic moiety of interest.

**[0028]** The displayed domain of a membrane fusion protein may be a binding moiety. By way of non-limiting example, binding moieties used for particular purposes may be a binding moiety directed to a compound or moiety displayed by a specific cell type or cells found predominantly in one type of tissue, which may be used to target minicells and their contents to specific cell types or tissues; or a binding moiety that is directed to a compound or moiety displayed by a pathogen, which may be used in diagnostic or therapeutic methods; a binding moiety that is directed to an undesirable compound, such as a toxin, which may be used to bind and preferably internalize and/or neutralize the undesirable compound; a diseased cell; or the binding moiety may be a domain that allows for the minicells to be covalently or non-covalently attached to a support material, which may be used in compositions and methods for compound screening and drug discovery. By “diseased cell” it is meant pathogen-infected cells, malfunctioning cells, and dysfunctional cells, e.g., cancer cells.

**[0029]** In various aspects, the minicells of the invention comprise one or more biologically active compounds. The term “biologically active” (synonymous with “bioactive”) indicates that a composition or compound itself has a biological effect, or that it modifies, causes, promotes, enhances, blocks, reduces, limits the production or activity of, or reacts with or binds to an endogenous molecule that has a biological effect. A “biological effect” may be but is not limited to one that stimulates or causes an immunoreactive response; one that impacts a biological process in an animal; one that impacts a biological process in a pathogen or parasite; one that generates or causes to be generated a detectable signal; and the like. Biologically active compositions, complexes or compounds may be used in therapeutic, prophylactic and diagnostic methods and compositions. Biologically active compositions, complexes or compounds act to cause or stimulate a desired effect upon an animal. Non-limiting examples of desired effects include, for example, preventing, treating or curing a disease or condition in an animal suffering therefrom; limiting the growth of or killing a pathogen in an animal infected thereby; augmenting the phenotype or genotype of an animal; stimulating a prophylactic immunoreactive response in an animal; or diagnosing a disease or disorder in an animal.

**[0030]** In the context of therapeutic applications of the invention, the term “biologically active” indicates that the composition, complex or compound has an activity that impacts an animal suffering from a disease or disorder in a positive sense and/or impacts a pathogen or parasite in a

negative sense. Thus, a biologically active composition, complex or compound may cause or promote a biological or biochemical activity within an animal that is detrimental to the growth and/or maintenance of a pathogen or parasite; or of cells, tissues or organs of an animal that have abnormal growth or biochemical characteristics, such as cancer cells.

[0031] In the context of diagnostic applications of the invention, the term "biologically active" indicates that the composition, complex or compound can be used for in vivo or ex vivo diagnostic methods and in diagnostic compositions and kits. For diagnostic purposes, a preferred biologically active composition or compound is one that can be detected, typically (but not necessarily) by virtue of comprising a detectable polypeptide. Antibodies to an epitope found on composition or compound may also be used for its detection.

[0032] In the context of prophylactic applications of the invention, the term "biologically active" indicates that the composition or compound induces or stimulates an immunoreactive response. In some preferred embodiments, the immunoreactive response is designed to be prophylactic, i.e., prevents infection by a pathogen. In other preferred embodiments, the immunoreactive response is designed to cause the immune system of an animal to react to the detriment of cells of an animal, such as cancer cells, that have abnormal growth or biochemical characteristics. In this application of the invention, compositions, complexes or compounds comprising antigens are formulated as a vaccine.

[0033] It will be understood by those skilled in the art that a given composition, complex or compound may be biologically active in therapeutic, diagnostic and prophylactic applications. A composition, complex or compound that is described as being "biologically active in a cell" is one that has biological activity in vitro (i.e., in a cell culture) or in vivo (i.e., in the cells of an animal). A "biologically active component" of a composition or compound is a portion thereof that is biologically active once it is liberated from the composition or compound. It should be noted, however, that such a component may also be biologically active in the context of the composition or compound.

[0034] In one aspect, the minicells of the invention comprise a therapeutic agent. Such minicells may be used to deliver therapeutic agents. In a preferred embodiment, a minicell comprising a therapeutic agent displays a binding moiety that specifically binds a ligand present on the surface of a cell, so that the minicells may be "targeted" to the cell. The therapeutic agent may be any type of compound or moiety, including without limitation small molecules, polypeptides, antibodies and antibody derivatives and nucleic acids. The therapeutic agent may be a drug; a prodrug, i.e., a compound that becomes biologically active in vivo after being introduced into a subject in need of treatment; or an immunogen.

[0035] In one aspect, the minicells of the invention comprise a detectable compound or moiety. As is understood by those of skill in the art, a compound or moiety that is "detectable" produces a signal that can be detected by spectroscopic, photochemical, biochemical, immunochemical, electromagnetic, radiochemical, or chemical means such as fluorescence, chemifluorescence, or chemiluminescence, electrochemiluminescence, or any other appropriate means.

A detectable compound may be a detectable polypeptide, and such polypeptides may, but need not, be incorporated into fusion membrane proteins of the minicell. Detectable polypeptides or amino acid sequences, includes, by way of non-limiting example, a green fluorescent protein (GFP), a luciferase, a beta-galactosidase, a His tag, an epitope, or a biotin-binding protein such as streptavidin or avidin. The detectable compound or moiety may be a radiolabeled compound or a radioisotope. A detectable compound or moiety may be a small molecule such as, by way of non-limiting example, a fluorescent dye; a radioactive isotope; or a compound that may be detected by x-rays or electromagnetic radiation. Image enhancers as those used for CAT and PET scans (e.g., calcium, gallidium) may be used. In another non-limiting example, detectable labels may also include loss of catalytic substrate or gain of catalytic product following catalysis by a minicell displayed, soluble cytoplasmic, or secreted enzyme.

[0036] In one aspect, the invention is drawn to a minicell comprising one or more bioactive nucleic acids or templates thereof. By way of non-limiting example, a bioactive nucleic acid may be an antisense oligonucleotide, an aptamer, an antisense transcript, a ribosomal RNA (rRNA), a transfer RNA (tRNA), a molecular decoy, or an enzymatically active nucleic acid, such as a ribozyme. Such minicells can, but need not, comprise a displayed polypeptide or protein on the surface of the minicell. The displayed polypeptide or protein may be a binding moiety directed to a compound or moiety displayed by a particular type of cell, or to a compound or moiety displayed by a pathogen. Such minicells can further, but need not, comprise an expression element having eubacterial, archaeal, eucaryotic, or viral expression sequences operably linked to a nucleotide sequence that serves as a template for a bioactive nucleic acid.

[0037] In one aspect, the invention is drawn to immunogenic minicells, i.e., minicells that display an immunogen, vaccines comprising immunogenic minicells, antibodies and antibody derivatives directed to immunogens displayed on immunogenic minicells, and method of making and using immunogenic minicells and antibodies and antibody derivatives produced therefrom in prophylactic, diagnostic, therapeutic and research applications. A preferred immunogen displayed by a minicell is an immunogenic polypeptide, which is preferably expressed from an expression element contained within the minicell in order to maximize the amount of immunogen displayed by the immunogenic minicells. The immunogenic polypeptide can be derived from any organism, obligate intracellular parasite, organelle or virus with the proviso that, in prophylactic applications, the immunogenic polypeptide is not derived from a prokaryote, including a eubacterial virus. The source organism for the immunogen may be a pathogen. A minicell displaying an immunogen derived from a pathogen is formulated into a vaccine and, in a prophylactic application, used to treat or prevent diseases and disorders caused by or related to the eukaryotic or archeobacterial pathogen.

[0038] In a separate aspect, the invention is drawn to minicells that display an immunogen derived from a non-functional, dysfunctional and/or diseased cell. By way of non-limiting example, the minicells display an immunogenic polypeptide derived from a hyperproliferative cell, i.e., a cell that is tumorigenic, or part of a tumor or cancer. As another non-limiting example, a cell that is infected with

a virus or an obligate intracellular parasite (e.g., Rickettsiae) displays an immunogenic polypeptide that is encoded by the genome of the infected cell but is aberrantly expressed in an infected cell. A vaccine comprising a minicell displaying an immunogen derived from a nonfunctional, dysfunctional and/or diseased cell is used in methods of treating or preventing hyperproliferative diseases or disorders, including without limitation a cell comprising an intracellular pathogen.

**[0039]** In one aspect, the invention is drawn to methods of using minicells, and expression systems optimized therefore, to manufacture, on a large scale, proteins using recombinant DNA technology. In a related aspect, the invention is drawn to the production, via recombinant DNA technology, and/or segregation of exogenous proteins in minicells. The minicells are enriched for the exogenous protein, which is desirable for increased yield and purity of the protein. In addition to protein purification, the minicells can be used for crystallography, the study of intracellular or extracellular protein-protein interactions, the study of intracellular or extracellular protein-nucleic acid interactions, the study of intracellular or extracellular protein-membrane interactions, and the study of other biological, chemical, or physiological event(s).

**[0040]** In one aspect, the invention is drawn to minicells having a membrane protein that has an intracellular domain. By way of non-limiting example, the intracellular domain is exposed on the inner surface of the minicell membrane oriented towards the cytoplasmic compartment. The intracellular protein domain is available for interaction with intracellular components. Intracellular components may be naturally present in the minicells or their parent cells, or may be introduced into minicells after segregation from parent cells. A membrane-associated protein may have more than one intracellular domain, and a minicell of the invention may display more than one membrane-associated protein.

**[0041]** In one aspect, the invention is drawn to a minicell comprising a membrane protein that is linked to a conjugatable compound (a.k.a. "attachable compound"). The conjugatable compound may be of any chemical nature and have one or more therapeutic or detectable moieties. By way of non-limiting example, a protein having a transmembrane or membrane anchoring domain is displayed and has the capacity to be specifically cross-linked on its extracellular domain. Through this approach, any conjugatable compound of interest may be quickly and easily attached to the outer surface of minicells containing this expressed membrane-spanning domain. In aspects of the invention wherein minicells are used for drug delivery *in vivo*, a preferred conjugatable compound is polyethylene glycol (PEG), which provides for "stealth" minicells that are not taken as well and/or as quickly by the reticuloendothelial system (RES). Other conjugatable compounds include polysaccharides, polynucleotides, lipopolysaccharides, lipoproteins, glycosylated proteins, synthetic chemical compounds, and/or chimeric combinations of these examples listed.

**[0042]** In various aspects of the invention, the minicell displays a polypeptide or other compound or moiety on its surface. By way of non-limiting example, a non-eubacterial membrane protein displayed by eubacterial minicells may be a receptor. Minicells displaying a receptor may, but need not, bind ligands of the receptor. In therapeutic applications of

this aspect of the invention, the ligand is an undesirable compound that is bound to its receptor and, in some aspects, is internalized by the minicells. In drug discovery applications of this aspect of the invention, the ligand for the receptor may be detectably labeled so that its binding to its receptor may be quantified. In the latter circumstance, the minicells may be used to identify and isolate, from a pool of compounds, one or more compounds that inhibit or stimulate the activity of the receptor. That is, these minicells can be used in screening assays, including assays such as those used in high throughput screening (HTS) systems and other drug discovery methods, for the purpose of preparing compounds that influence the activity of a receptor of interest.

**[0043]** The non-eubacterial membrane protein displayed by minicells may be a fusion protein, i.e., a protein that comprises a first polypeptide having a first amino acid sequence and a second polypeptide having a second amino acid sequence, wherein the first and second amino acid sequences are not naturally present in the same polypeptide. At least one polypeptide in a membrane fusion protein is a "transmembrane domain" or "membrane-anchoring domain". The transmembrane and membrane-anchoring domains of a membrane fusion protein may be selected from membrane proteins that naturally occur in a eukaryote, such as a fungus, a unicellular eukaryote, a plant and an animal, such as a mammal including a human. Such domains may be from a viral membrane protein naturally found in a virus such as a bacteriophage or a eukaryotic virus, e.g., an adenovirus or a retrovirus. Such domains may be from a membrane protein naturally found in an archaeobacterium such as a thermophile.

**[0044]** The displayed domain of a membrane fusion protein may be an enzymatic domain such as one having the activity of a lipase, a kinase, a phosphatase, a reductase, a protease, or a nuclease. Contacting such minicells with the appropriate substrate of the enzyme allows the substrate to be converted to reactant. When either the substrate or reactant is detectable, the reaction catalyzed by the membrane-bound enzyme may be quantified. In the latter instance, the minicells may be used to identify and isolate, from a pool of compounds, one or more compounds that inhibit or stimulate the activity of the enzyme represented by the displayed enzymatic moiety. That is, these minicells can be used in screening assays, including assays such as those used in high throughput screening (HTS) systems and other drug discovery methods, for the purpose of preparing compounds that influence the activity of an enzyme or enzymatic moiety of interest.

**[0045]** The displayed domain of a membrane fusion protein may be a binding moiety. By way of non-limiting example, binding moieties used for particular purposes may be a binding moiety directed to a compound or moiety displayed by a specific cell type or cells found predominantly in one type of tissue, which may be used to target minicells and their contents to specific cell types or tissues; or a binding moiety that is directed to a compound or moiety displayed by a pathogen, which may be used in diagnostic or therapeutic methods; a binding moiety that is directed to an undesirable compound, such as a toxin, which may be used to bind and preferably internalize and/or neutralize the undesirable compound; a diseased cell; or the binding moiety may be a domain that allows for the minicells to be covalently or non-covalently attached to a support material,

which may be used in compositions and methods for compound screening and drug discovery.

**[0046]** In one aspect, the invention provides compositions and methods for preparing a soluble and/or secreted protein where the protein remains in the cytoplasm of the minicell or is secreted following native secretory pathways for endogenous secreted proteins or is secreted using chimeric fusion to secretory signaling sequences. By way of non-limiting example, secreted or cytoplasmic soluble proteins may be produced for purification, targeted therapeutic applications where the protein produced is a therapeutic agent and is produced at the desired site of, detection for screening or diagnostic purposes where the protein is produced in response to a stimulus and/or localization event, or to stimulate targeted minicell-cell fusion or interaction events where the protein produced stimulates cell-cell fusion upon targeted stimulation.

**[0047]** In one aspect, the invention provides compositions and methods for preparing antibodies and/or antibody derivatives that recognize an immunogenic epitope present on the native form of a membrane protein, but which is not immunogenic when the membrane protein is denatured or when prepared as a synthetic oligopeptide. Such antibodies and antibody derivatives are said to be "conformation sensitive." Unlike most antibodies and antibody derivatives prepared by using a denatured membrane protein or an oligopeptide derived from the membrane protein, conformation sensitive antibodies and antibody derivatives specifically bind membrane proteins in their native state (i.e., in a membrane) with high affinity. Conformation sensitive antibodies and antibody derivatives are used to target compounds and compositions, including a minicell of the invention, to a cell displaying the membrane protein of choice. Conformation sensitive antibodies and antibody derivatives are also used to prevent receptors from binding their natural ligands by specifically binding to the receptor with a high affinity and thereby limiting access of the ligand to the receptor. Conformation sensitive antibodies and antibody derivatives can be prepared that are specific for a specific isoform or mutant of a membrane protein, which can be useful in research and medical applications.

**[0048]** In one aspect, the invention provides biosensors comprising minicells including, not limited to, the minicells of the invention. An exemplary biosensor of the invention is a BIAcore chip, i.e., a chip onto which minicells are attached, where the minicells undergo some change upon exposure to a preselected compound, and the change is detected using surface plasmon resonance. A biosensor comprising minicells can be used in methods of detecting the presence of an undesirable compound. Undesirable compounds include but are not limited to, toxins; pollutants; explosives, such as those in landmines or illegally present; illegal narcotics; components of biological or chemical weapons. In a related aspect, the invention provides a device comprising a microchip operatively associated with a biosensor comprising a minicell. The device can further comprise an actuator that performs a responsive function when the sensor detects a preselected level of a marker.

**[0049]** In one aspect, the invention provides minicells that may be used as research tools and/or kits comprising such research tools. The minicells of the invention may be used as is, or incorporated into research tools useful for scientific

research regarding all amino acid comprising compounds including, but not limited to membrane-associated proteins, chimeric membrane fusion proteins, and soluble proteins. Such scientific research includes, by way of non-limiting example, basic research, as well as pharmacological, diagnostic, and pharmacogenetic studies. Such studies may be carried out in vivo or in vitro.

**[0050]** In one aspect, the invention is drawn to archaeobacterial minicells. In a related aspect, the invention is drawn to archaeobacterial minicells comprising at least one exogenous protein, that is, a protein that is not normally found in the parent cell, including without limitation fusion proteins. The archaeobacterial minicells of the invention optionally comprise an expression element that directs the production of the exogenous protein(s).

**[0051]** In other aspects, the invention is drawn to methods of preparing the minicells, protoplasts, and poroplasts<sup>TM</sup> of the invention for various applications including but not limited to diagnostic, therapeutic, research and screening applications. In a related aspect, the invention is drawn to pharmaceutical compositions, reagents and kits comprising minicells.

**[0052]** In each aspect and embodiment of the invention, unless stated otherwise, embodiments wherein the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast exist.

**[0053]** In a first aspect, the invention provides a minicell comprising a membrane protein selected from the group consisting of a eukaryotic membrane protein, an archaeobacterial membrane protein and an organellar membrane protein. In another embodiment, wherein the minicell comprises a biologically active compound. By way of non-limiting example, the biologically active compound is a radioisotope, a polypeptide, a nucleic acid or a small molecule.

**[0054]** In another embodiment, the minicell comprises an expression construct, wherein the first expression construct comprises expression sequences operably linked to an ORF that encodes a protein. In another embodiment, the ORF encodes the membrane protein. In another embodiment, the expression sequences that are operably linked to an ORF are inducible and/or repressible.

**[0055]** In another aspect, the minicell comprises a second expression construct, wherein the second expression construct comprises expression sequences operably linked to a gene. In another embodiment, the expression sequences that are operably linked to a gene are inducible and/or repressible. In a related embodiment, the gene product of the gene regulates the expression of the ORF that encodes the protein. A factor that "regulates" the expression of a gene or a gene product directly or indirectly initiates, enhances, quickens, slows, terminates, limits or completely blocks expression of a gene. In different embodiments, the gene product of the gene is a nucleic acid or a polypeptide. The polypeptide can be of any type, including but not limited to a membrane protein, a soluble protein or a secreted protein. A membrane protein can be a membrane fusion protein comprising a first polypeptide, which comprises at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide.

**[0056]** In one aspect, the invention provides a minicell comprising a membrane fusion protein, the fusion protein

comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide, wherein the second polypeptide is not derived from a eubacterial protein and is neither a His tag nor a glutathione-S-transferase polypeptide. In various embodiments, the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast. In one embodiment, the minicell comprises a biologically active compound.

[0057] In one aspect, the invention provides a minicell comprising a membrane conjugate, wherein the membrane conjugate comprises a membrane protein chemically linked to a conjugated compound. In one embodiment, the conjugated compound is selected from the group consisting of a nucleic acid, a polypeptide, a lipid and a small molecule.

[0058] In one aspect, the invention provides a method for making minicells, comprising (a) culturing a minicell-producing parent cell, wherein the parent cell comprises an expression construct, wherein the expression construct comprises a gene operably linked to expression sequences that are inducible and/or repressible, and wherein induction or repression of the gene causes or enhances the production of minicells; and (b) separating the minicells from the parent cell, thereby generating a composition comprising minicells, wherein an inducer or repressor is present within the parent cells during one or more steps and/or between two or more steps of the method. In one embodiment, the method further comprises (c) purifying the minicells from the composition.

[0059] Relevant gene products are factors involved in or modulating DNA replication, cellular division, cellular partitioning, septation, transcription, translation, or protein folding. The minicells are separated from parent cells by processes such as centrifugation, ultracentrifugation, density gradation, immunoaffinity, immunoprecipitation and other techniques described herein.

[0060] In one embodiment, the minicell is a poroplast, and the method further comprises (d) treating the minicells with an agent, or incubating the minicells under a set of conditions, that degrades the outer membrane of the minicell. The outer membrane is degraded by treatment with an agent selected from the group consisting of EDTA, EGTA, lactic acid, citric acid, gluconic acid, tartaric acid, polyethyleneimine, polycationic peptides, cationic leukocyte peptides, aminoglycosides, aminoglycosides, protamine, insect cecropins, reptilian magainins, polymers of basic amino acids, polymixin B, chloroform, nitrilotriacetic acid and sodium hexametaphosphate; by exposure to conditions selected from the group consisting of osmotic shock and insonation; and by other methods described herein.

[0061] In one embodiment, further comprising removing one or more contaminants from the composition. Representative contaminants are LPS and peptidoglycan. In a representative embodiment, LPS is removed by contacting the composition to an agent that binds or degrades LPS. At least about 50%, preferably about 65% to about 75%, more preferably 95%, most preferably 99% or >99% of LPS is removed from an initial preparation of minicells. In a related embodiment, the minicell-producing parent cell comprises a mutation in a gene required for lipopolysaccharide synthesis.

[0062] In one embodiment, the minicell is a spheroplast, and the method further comprises (d) treating the minicells

with an agent, or incubating the minicells under a set of conditions, that disrupts or degrades the outer membrane; and (e) treating the minicells with an agent, or incubating the minicells under a set of conditions, that disrupts or degrades the cell wall. The agent that disrupts or degrades the cell wall can be, e.g., a lysozyme, and the set of conditions that disrupts or degrades the cell wall can be, e.g., incubation in a hypertonic solution.

[0063] In one embodiment, the minicell is a protoplast, and the method further comprises (d) treating the minicells with an agent, or incubating the minicells under a set of conditions, that disrupt or degrades the outer membrane; (e) treating the minicells with an agent, or incubating the minicells under a set of conditions, that disrupts or degrades the cell wall, in order to generate a composition that comprises protoplasts; and (f) purifying protoplasts from the composition. In one embodiment, the method further comprises preparing a denuded minicell from the minicell. In one embodiment, the method further comprises covalently or non-covalently linking one or more components of the minicell to a conjugated moiety.

[0064] In one aspect, the invention provides a L-form minicell comprising (a) culturing an L-form eubacterium, wherein the eubacterium comprises one or more of the following: (i) an expression element that comprises a gene operably linked to expression sequences that are inducible and/or repressible, wherein induction or repression of the gene regulates the copy number of an episomal expression construct; (ii) a mutation in an endogenous gene, wherein the mutation regulates the copy number of an episomal expression construct; (iii) an expression element that comprises a gene operably linked to expression sequences that are inducible and/or repressible, wherein induction or repression of the gene causes or enhances the production of minicells; and (iv) a mutation in an endogenous gene, wherein the mutation causes or enhances minicell production; (b) culturing the L-form minicell-producing parent cell in media under conditions wherein minicells are produced; and (c) separating the minicells from the parent cell, thereby generating a composition comprising L-form minicells, wherein an inducer or repressor is present within the minicells during one or more steps and/or between two or more steps of the method. In one embodiment, the method further comprises (d) purifying the L-form minicells from the composition.

[0065] In one aspect, the invention provides a method of producing a protein, comprising (a) transforming a minicell-producing parent cell with an expression element that comprises expression sequences operably linked to a nucleic acid having an ORF that encodes the protein; (b) culturing the minicell-producing parent cell under conditions wherein minicells are produced; and (c) purifying minicells from the parent cell, (d) purifying the protein from the minicells, wherein the ORF is expressed during step (b), between steps (b) and (c), and during step (c).

[0066] In one embodiment, the expression elements segregate into the minicells, and the ORF is expressed between steps (c) and (d). In one embodiment, the protein is a soluble protein contained within the minicells, and the method further comprises (e) lysing the minicells.

[0067] In one embodiment, the protein is a secreted protein, and the method further comprises (e) collecting a

composition in which the minicells are suspended or with which the minicells are in contact.

[0068] In one embodiment, the expression sequences to which the ORF is operably linked are inducible, wherein the method further comprises adding an inducing agent between steps (a) and (b); during step (b); and between steps (b) and (c).

[0069] In one embodiment, the expression sequences to which the ORF is operably linked are inducible, wherein the expression elements segregate into the minicells, the method further comprises adding an inducing agent after step (c).

[0070] In one embodiment, the method further comprises (e) preparing poroplasts from the minicells, wherein the ORF is expressed during step (b); between steps (b) and (c); during step (c); between step (c) and step (d) when the expression elements segregate into the minicells; and/or after step (d) when the expression elements segregate into the minicells.

[0071] In one embodiment, the method further comprises (f) purifying the protein from the poroplasts.

[0072] In one embodiment, the method further comprises (e) preparing spheroplasts from the minicells, wherein the ORF is expressed during step (b), between steps (b) and (c), during step (c), between steps (c) and (d) and/or after step (d).

[0073] In one embodiment, the method further comprises (f) purifying the protein from the spheroplasts.

[0074] In one embodiment, the method further comprises (e) preparing protoplasts from the minicells, wherein the ORF is expressed during step (b), between steps (b) and (c), during step (c), between steps (c) and (d) and/or after step (d).

[0075] In one embodiment, the method further comprises (f) purifying the protein from the protoplasts.

[0076] In one embodiment, the method further comprises (e) preparing membrane preparations from the minicells, wherein the ORF is expressed during step (b), between steps (b) and (c), during step (c), between steps (c) and (d) and/or after step (d).

[0077] In one embodiment, the method further comprises (f) purifying the protein from the membrane preparations.

[0078] In one embodiment, the minicell-producing parent cell is an L-form bacterium.

[0079] In one aspect, the invention provides a method of producing a protein, comprising (a) transforming a minicell with an expression element that comprises expression sequences operably linked to a nucleic acid having an ORF that encodes the protein; and (b) incubating the minicells under conditions wherein the ORF is expressed.

[0080] In one embodiment, the method further comprises (c) purifying the protein from the minicells.

[0081] In one aspect, the invention provides a method of producing a protein, comprising (a) transforming a minicell-producing parent cell with an expression element that comprises expression sequences operably linked to a nucleic acid having an ORF that encodes a fusion protein comprising the protein and a polypeptide, wherein a protease-

sensitive amino acid sequence is positioned between the protein and the polypeptide; (b) culturing the minicell-producing parent cell under conditions wherein minicells are produced; (c) purifying minicells from the parent cell, wherein the ORF is expressed during step (b); between steps (b) and (c); and/or after step (c) when the expression elements segregate into the minicells; and (d) treating the minicells with a protease that cleaves the sensitive amino acid sequence, thereby separating the protein from the polypeptide.

[0082] In one aspect, the invention provides a poroplast, the poroplast comprising a vesicle, bonded by a membrane, wherein the membrane is an eubacterial inner membrane, wherein the vesicle is surrounded by a eubacterial cell wall, and wherein the eubacterial inner membrane is accessible to a compound in solution with the poroplast. In one embodiment, the poroplast is a cellular poroplast. The compound has a molecular weight of at least 1 kD, preferably at least about 0.1 to about 1 kD, more preferably from about 1, 10 or 25 kD to about 50 kD, and most preferably from about 75 or about 100 kD to about 150 or 300 kD.

[0083] In one embodiment, the poroplast comprises an exogenous nucleic acid, which may be an expression construct. In one embodiment, the expression construct comprises an ORF that encodes an exogenous protein, wherein the ORF is operably linked to expression sequences. In one embodiment, the exogenous protein is a fusion protein, a soluble protein or a secreted protein. In one embodiment, the exogenous protein is a membrane protein, and is preferably accessible to compounds in solution with the poroplast. In one embodiment, poroplasts are placed in a hypertonic solution, wherein 90% or more of an equivalent amount of spheroplasts or protoplasts lyse in the solution under the same conditions.

[0084] In one embodiment, the membrane protein is selected from the group consisting of a eukaryotic membrane protein, an archaeobacterial membrane protein, and an organellar membrane protein. In one embodiment, the membrane protein is a fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide, wherein the second polypeptide is displayed by the poroplast. In one embodiment, the second polypeptide is displayed on the external side of the eubacterial inner membrane. The second polypeptide can be an enzyme moiety, a binding moiety, a toxin, a cellular uptake sequence, an epitope, a detectable polypeptide, and a polypeptide comprising a conjugatable moiety. An enzyme moiety is a polypeptide derived from, by way of non-limiting example, a cytochrome P450, an oxidoreductase, a transferase, a hydrolase, a lyase, an isomerase, a ligase or a synthetase.

[0085] In one embodiment, the poroplast comprises a membrane component that is chemically linked to a conjugated compound.

[0086] In one embodiment, the expression construct comprises one or more DNA fragments from a genome or cDNA. In one embodiment, the exogenous protein has a primary amino acid sequence predicted from a nucleic acid sequence.

[0087] In one aspect, the invention provides a solid support comprising a minicell. In various embodiments, the

solid support is a dipstick, a bead or a microtiter multiwell plate. In one embodiment, the minicell comprises a detectable compound, which may be a colorimetric, fluorescent or radioactive compound.

[0088] In one embodiment, the minicell displays a membrane component selected from the group consisting of (i) a eukaryotic membrane protein, (ii) an archeobacterial membrane protein, (iii) an organellar membrane protein, (iv) a fusion protein comprising at least one transmembrane domain or at least one membrane anchoring domain, and (v) a membrane conjugate comprising a membrane component chemically linked to a conjugated compound.

[0089] In one embodiment, the membrane component is a receptor. In a related embodiment, the solid support further comprises a co-receptor. In one embodiment, the minicell displays a binding moiety.

[0090] In one aspect, the invention provides a solid support comprising a minicell, wherein the minicell displays a fusion protein, the fusion protein comprising a first polypeptide that comprises at least one transmembrane domain or at least one membrane anchoring domain, and a second polypeptide. In various embodiments, the second polypeptide comprises a binding moiety or an enzyme moiety.

[0091] In one aspect, the invention provides a solid support comprising a minicell, wherein the minicell comprises a membrane conjugate comprising a membrane component chemically linked to a conjugated compound. In one embodiment, the conjugated compound is a spacer. In one embodiment, the spacer is covalently linked to the solid support. In one embodiment, the conjugated compound is covalently linked to the solid support.

[0092] In one aspect, the invention provides a minicell comprising a biologically active compound, wherein the minicell displays a ligand or binding moiety, wherein the ligand or binding moiety is part of a fusion protein comprising a first polypeptide that comprises at least one transmembrane domain or at least one membrane anchoring domain and a second polypeptide that comprises a binding moiety, and the minicell is a poroplast, spheroplast or protoplast.

[0093] In one aspect, the invention provides a eubacterial minicell comprising a biologically active compound, wherein the minicell displays a binding moiety, wherein the binding moiety is selected from the group consisting of (a) a eukaryotic membrane protein; (b) an archeobacterial membrane protein; (c) an organellar membrane protein; and (d) a fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide, wherein the second polypeptide is not derived from a eubacterial protein and is neither a His tag nor a glutathione-S-transferase polypeptide, and wherein the polypeptide comprises a binding moiety.

[0094] In one embodiment, the binding moiety is selected from the group consisting of an antibody, an antibody derivative, a receptor and an active site of a non-catalytic derivative of an enzyme. In a preferred embodiment, the binding moiety is a single-chain antibody. In one embodiment, one of the ORFs encodes a protein that comprises the binding moiety.

[0095] In one embodiment, the binding moiety is directed to a ligand selected from the group consisting of an epitope displayed on a pathogen, an epitope displayed on an infected cell and an epitope displayed on a hyperproliferative cell.

[0096] In one embodiment, the invention further comprises a first and second nucleic acid, wherein the first nucleic acid comprises eukaryotic expression sequences operably linked to a first ORF, and a second nucleic acid, wherein the second nucleic acid comprises eubacterial expression sequences operably linked to a second ORF.

[0097] In one embodiment, the eubacterial expression sequences are induced and/or derepressed when the binding moiety is in contact with a target cell. In a variant embodiment, the eukaryotic expression sequences are induced and/or derepressed when the nucleic acid is in the cytoplasm of a eukaryotic cell. In related embodiments, the protein encoded by the first ORF comprises eukaryotic secretion sequences and/or the protein encoded by the second ORF comprises eubacterial secretion sequences.

[0098] In one aspect, the invention provides a method of associating a radioactive compound with a cell, wherein the cell displays a ligand specifically recognized by a binding moiety, comprising contacting the cell with a minicell that comprises the radioactive compound and displays the binding moiety. In a diagnostic embodiment, the amount of radiation emitted by the radioactive isotope is sufficient to be detectable. In a therapeutic embodiment, the amount of radiation emitted by the radioactive isotope is sufficient to be cytotoxic. In one embodiment, the ligand displayed by the cell is selected from the group consisting of an epitope displayed on a pathogen, an epitope displayed on an infected cell and an epitope displayed on a hyperproliferative cell. In one embodiment, the binding moiety is selected from the group consisting of an antibody, an antibody derivative, a channel protein and a receptor, and is preferably a single-chain antibody. In other embodiments, the binding moiety is an aptamer or a small molecule. In one embodiment, the ligand is selected from the group consisting of a cytokine, hormone, and a small molecule.

[0099] In one aspect, the invention provides a method of delivering a biologically active compound to a cell, wherein the cell displays a ligand specifically recognized by a binding moiety, comprising contacting the cell with a minicell that displays the binding moiety, wherein the minicell comprises the biologically active compound, and wherein the contents of the minicell are delivered into the cell from a minicell bound to the cell. In one embodiment, the biologically active compound is selected from the group consisting of a nucleic acid, a lipid, a polypeptide, a radioactive compound, an ion and a small molecule.

[0100] In one embodiment, the membrane of the minicell comprises a system for transferring a molecule from the interior of a minicell into the cytoplasm of the cell. A representative system for transferring a molecule from the interior of a minicell into the cytoplasm of the cell is a Type III secretion system.

[0101] In one embodiment, the minicell further comprises a first and second nucleic acid, wherein the first nucleic acid comprises eukaryotic expression sequences operably linked to a first ORF, and a second nucleic acid, wherein the second nucleic acid comprises eubacterial expression sequences



operably linked to a second ORF. In one embodiment, one of the ORFs encodes a protein that comprises the binding moiety. In one embodiment, the eubacterial expression sequences are induced and/or derepressed when the binding moiety is in contact with a target cell. In one embodiment, the eukaryotic expression sequences are induced and/or derepressed when the nucleic acid is in the cytoplasm of a eukaryotic cell. In one embodiment, the protein encoded by the first ORF comprises eukaryotic secretion sequences and/or the protein encoded by the second ORF comprises eubacterial secretion sequences. In one embodiment, the ligand is selected from the group consisting of a cytokine, hormone, and a small molecule.

[0102] In one aspect, the invention provides a minicell displaying a synthetic linking moiety, wherein the synthetic linking moiety is covalently or non-covalently attached to a membrane component of the minicell.

[0103] In one aspect, the invention provides a sterically stabilized minicell comprising a displayed moiety that has a longer half-life in vivo than a wild-type minicell, wherein the displayed moiety is a hydrophilic polymer that comprises a PEG moiety, a carboxylic group of a polyalkylene glycol or PEG stearate.

[0104] In one aspect, the invention provides a minicell having a membrane comprising an exogenous lipid, wherein a minicell comprising the exogenous lipid has a longer half-life in vivo than a minicell lacking the exogenous lipid, and wherein the minicell is selected from the group consisting of a eubacterial minicell, a poroplast, a spheroplast and a protoplast. In one embodiment, the exogenous lipid is a derivitized lipid which may, by way of non-limiting example, be phosphatidylethanolamine derivitized with PEG, DSPE-PEG, PEG stearate; PEG-derivitized phospholipids, a PEG ceramide or DSPE-PEG.

[0105] In one embodiment, the exogenous lipid is not present in a wild-type membrane, or is present in a different proportion than is found in minicells comprising a wild-type membrane. The exogenous lipid can be a ganglioside, sphingomyelin, monosialoganglioside GM1, galactocerebroside sulfate, 1,2-sn-dimyristoylphosphatidylcholine, phosphatidylinositol and cardiolipin.

[0106] In one embodiment, the linking moiety is non-covalently attached to the minicell. In one embodiment, one of the linking moiety and the membrane component comprises biotin, and the other comprises avidin or streptavidin. In one embodiment, the synthetic linking moiety is a cross-linker. In one embodiment, the cross-linker is a bifunctional cross-linker.

[0107] In one aspect, the invention provides a method of transferring a membrane protein from a minicell membrane to a biological membrane comprising contacting a minicell to the biological membrane, wherein the minicell membrane comprises the membrane protein, and allowing the minicell and the biological membrane to remain in contact for a period of time sufficient for the transfer to occur.

[0108] In one embodiment, the biological membrane is a cytoplasmic membrane or an organellar membrane. In one embodiment, the biological membrane is a membrane selected from the group consisting of a membrane of a pathogen, a membrane of an infected cell and a membrane of a hyperproliferative cell. In one embodiment, the biological

membrane is the cytoplasmic membrane of a recipient cell, which may be a cultured cell and a cell within an organism. In one embodiment, the biological membrane is present on a cell that has been removed from an animal, the contacting occurs in vitro, after which the cell is returned to the organism.

[0109] In one embodiment, the membrane protein is an enzyme. In this embodiment, the membrane protein having enzymatic activity is selected from the group consisting of a cytochrome P450 and a fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one polypeptide, wherein the second polypeptide has enzymatic activity.

[0110] In one embodiment, the membrane protein is a membrane fusion protein, the membrane fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide.

[0111] In one embodiment, the second polypeptide is a biologically active polypeptide. In one embodiment, the minicell displays ligand or a binding moiety.

[0112] In one aspect, the invention provides a minicell that comprises an expression construct comprising an ORF encoding a membrane protein operably linked to expression sequences, wherein the expression sequences are induced and/or derepressed when the minicell is in contact with a target cell.

[0113] In one embodiment, the biological membrane is a cytoplasmic membrane or an organellar membrane. In one embodiment, the biological membrane is a membrane selected from the group consisting of a membrane of a pathogen, a membrane of an infected cell and a membrane of a hyperproliferative cell. In one embodiment, the minicell displays a ligand or a binding moiety selected from the group consisting of an antibody, an antibody derivative, an aptamer and a small molecule. In one embodiment, the membrane protein is a membrane fusion protein, the membrane fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide. In one embodiment, the ligand is selected from the group consisting of a cytokine, hormone, and a small molecule.

[0114] In one aspect, the invention provides a pharmaceutical composition comprising a minicell, wherein the minicell displays a membrane protein, wherein the membrane protein is selected from the group consisting of a eukaryotic membrane protein, an archeobacterial membrane protein and an organellar membrane protein. In one embodiment, the membrane protein is selected from the group consisting of a receptor, a channel protein, a cellular adhesion factor and an integrin. In one embodiment, the pharmaceutical formulation further comprises an adjuvant. In one embodiment, the membrane protein comprises a polypeptide epitope displayed by a hyperproliferative cell. In one embodiment, the membrane protein comprises an epitope displayed by a eukaryotic pathogen, an archeobacterial pathogen, a virus or an infected cell.

[0115] In one aspect, the invention provides a pharmaceutical composition comprising a minicell, wherein the minicell displays a membrane protein that is a fusion protein, the

fusion protein comprising (i) a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and (ii) a second polypeptide, wherein the second polypeptide is not derived from a eubacterial protein. In one embodiment, the pharmaceutical formulation further comprises an adjuvant. In one embodiment, the second polypeptide comprises a polypeptide epitope displayed by a hyperproliferative cell. In one embodiment, the membrane protein comprises an epitope displayed by a eukaryotic pathogen, an archeabacterial pathogen, a virus or an infected cell.

[0116] In one aspect, the invention provides a pharmaceutical composition comprising a minicell, wherein the minicell displays a membrane conjugate, wherein the membrane conjugate comprises a membrane component chemically linked to a conjugated compound. In one embodiment, the membrane protein is selected from the group consisting of a receptor, a channel protein, a cellular adhesion factor and an integrin. In one embodiment, the pharmaceutical further comprises an adjuvant. In one embodiment, the membrane component is a polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain, or a lipid that is part of a membrane. In one embodiment, the conjugated compound is a polypeptide, and the chemical linkage between the membrane component and the conjugated compound is not a peptide bond. In one embodiment, the conjugated compound is a nucleic acid. In one embodiment, the conjugated compound is an organic compound. In one embodiment, the organic compound is selected from the group consisting of a narcotic, a toxin, a venom, a sphingolipid and a soluble protein.

[0117] In one aspect, the invention provides a method of making a pharmaceutical composition comprising a minicell, wherein the minicell displays a membrane protein, wherein the membrane protein is selected from the group consisting of a eukaryotic membrane protein, an archeabacterial membrane protein and an organellar membrane protein. In one embodiment, the method further comprises adding an adjuvant to the pharmaceutical formulation. In one embodiment, the method further comprises desiccating the formulation. In one embodiment, the method further comprises adding a suspension buffer to the formulation. In one embodiment, the method further comprises making a chemical modification of the membrane protein. In one embodiment, the chemical modification is selected from the group consisting of glycosylation, deglycosylation, phosphorylation, dephosphorylation and proteolysis. In one aspect, the invention provides a method of making a pharmaceutical composition comprising a minicell, wherein the minicell displays a membrane protein that is a fusion protein, the fusion protein comprising (i) a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and (ii) a second polypeptide, wherein the second polypeptide is not derived from a eubacterial protein.

[0118] In one aspect, the invention provides a method of making a pharmaceutical formulation comprising a minicell, wherein the minicell displays a membrane conjugate, wherein the membrane conjugate comprises a membrane component chemically linked to a conjugated compound. In one embodiment, the method further comprises adding an adjuvant to the pharmaceutical formulation. In one embodiment, the membrane component is a polypeptide comprising

at least one transmembrane domain or at least one membrane anchoring domain, or a lipid that is part of a membrane. In one embodiment, the conjugated compound is a polypeptide, and the chemical linkage between the membrane component and the conjugated compound is not a peptide bond. In one embodiment, the conjugated compound is a nucleic acid. In one embodiment, the conjugated compound is an organic compound. In one embodiment, the organic compound is selected from the group consisting of a narcotic, a toxin, a venom, and a sphingolipid.

[0119] In one aspect, the invention provides a method of detecting an agent that is specifically bound by a binding moiety, comprising contacting a minicell displaying the binding moiety with a composition known or suspected to contain the agent, and detecting a signal that is modulated by the binding of the agent to the binding moiety. In one embodiment, the agent is associated with a disease. In one embodiment, the minicell comprises a detectable compound. In one embodiment, the binding moiety is antibody or antibody derivative. In one embodiment, the composition is an environmental sample. In one embodiment, the composition is a biological sample. In one embodiment, the biological sample is selected from the group consisting of blood, serum, plasma, urine, saliva, a biopsy sample, feces and a skin patch.

[0120] In one aspect, the invention provides a method of *in situ* imaging of a tissue or organ, comprising administering to an organism a minicell comprising an imaging agent and a binding moiety and detecting the imaging agent in the organism.

[0121] In one embodiment, the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast. In one embodiment, the binding moiety is an antibody or antibody derivative. In one embodiment, the binding moiety specifically binds a cell surface antigen. In one embodiment, the cell surface antigen is an antigen displayed by a tumorigenic cell, a cancer cell, and an infected cell. In one embodiment, the cell surface antigen is a tissue-specific antigen. In one embodiment, the method of imaging is selected from the group consisting of magnetic resonance imaging, ultrasound imaging; and computer axial tomography (CAT). In one aspect, the invention provides a device comprising a microchip operatively associated with a biosensor comprising a minicell, wherein the microchip comprises or contacts the minicell, and wherein the minicell displays a binding moiety.

[0122] In one embodiment, the invention provides a method of detecting a substance that is specifically bound by a binding moiety, comprising contacting the device of claim 16 with a composition known or suspected to contain the substance, and detecting a signal from the device, wherein the signal changes as a function of the amount of the substance present in the composition. In one embodiment, the composition is a biological sample or an environmental sample.

[0123] In one aspect, the invention provides a method of identifying an agent that specifically binds a target compound, comprising contacting a minicell displaying the target compound with a library of compounds, and identifying an agent in the library that binds the target compound. In one embodiment, the library of compounds is a protein library. In one embodiment, the protein library is selected

from the group consisting of a phage display library, a phagemid display library, a baculovirus library, a yeast display library, and a ribosomal display library. In one embodiment, the library of compounds is selected from the group consisting of a library of aptamers, a library of synthetic peptides and a library of small molecules.

[0124] In one embodiment, the target compound is a target polypeptide. In one embodiment, the minicell comprises an expression construct comprising expression sequences operably linked to an ORF encoding the target polypeptide. In one embodiment, the target polypeptide is a membrane protein. In one embodiment, the membrane protein is a receptor or a channel protein. In one embodiment, the membrane protein is an enzyme. In one embodiment, the target compound is a membrane fusion protein, the membrane fusion protein comprising a first polypeptide, wherein the first polypeptide comprises at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide, wherein the second polypeptide comprises amino acid sequences derived from a target polypeptide. In one embodiment, the method further comprises comparing the activity of the target compound in the presence of the agent to the activity of the target compound in the absence of the agent.

[0125] In one embodiment, the activity of the target compound is an enzyme activity. In one embodiment, the activity of the target compound is a binding activity. In one embodiment, the invention further comprises comparing the binding of the agent to the target compound to the binding of a known ligand of the target compound. In one embodiment, a competition assay is used for the comparing.

[0126] In one aspect, the invention provides a device comprising microchips operatively associated with a biosensor comprising a set of minicells in a prearranged pattern, wherein each of the microchips comprise or contact a minicell, wherein each of the minicell displays a different target compound, and wherein binding of a ligand to a target compound results in an increased or decreased signal. In one embodiment, the invention provides a method of identifying an agent that specifically binds a target compound, comprising contacting the device with a library of compounds, and detecting a signal from the device, wherein the signal changes as a function of the binding of an agent to the target compound. In one embodiment, the invention provides a method of identifying an agent that specifically blocks the binding of a target compound to its ligand, comprising contacting the device with a library of compounds, and detecting a signal from the device, wherein the signal changes as a function of the binding of an agent to the target compound.

[0127] In one aspect, the invention provides a method of making an antibody that specifically binds a protein domain, wherein the domain is in its native conformation, wherein the domain is contained within a protein displayed on a minicell, comprising contacting the minicell with a cell, wherein the cell is competent for producing antibodies to an antigen contacted with the cell, in order to generate an immunogenic response in which the cell produces the antibody.

[0128] In one embodiment, the protein displayed on a minicell is a membrane protein. In one embodiment, the membrane protein is a receptor or a channel protein. In one

embodiment, the domain is found within the second polypeptide of a membrane fusion protein, wherein the membrane fusion protein comprises a first polypeptide, wherein the first polypeptide comprises at least one transmembrane domain or at least one membrane anchoring domain. In one embodiment, the contacting occurs *in vivo*. In one embodiment, the antibody is a polyclonal antibody or a monoclonal antibody. In one embodiment, the contacting occurs in an animal that comprises an adjuvant.

[0129] In one aspect, the invention provides the method of making an antibody derivative that specifically binds a protein domain, wherein the domain is in its native conformation, wherein the domain is displayed on a minicell, comprising contacting the minicell with a protein library, and identifying an antibody derivative from the protein library that specifically binds the protein domain. In one embodiment, the protein library is selected from the group consisting of a phage display library, a phagemid display library, and a ribosomal display library.

[0130] In one aspect, the invention provides a method of making an antibody or antibody derivative that specifically binds an epitope, wherein the epitope is selected from the group consisting of (i) an epitope composed of amino acids found within a membrane protein, (ii) an epitope present in an interface between a membrane protein and a membrane component, (iii) an epitope present in an interface between a membrane protein and one or more other proteins and (iv) an epitope in a fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain, and a second polypeptide, the second polypeptide comprising the epitope; comprising contacting a minicell displaying the epitope with a protein library, or to a cell, wherein the cell is competent for producing antibodies to an antigen contacted with the cell, in order to generate an immunogenic response in which the cell produces the antibody.

[0131] In one embodiment, the cell is contacted *in vivo*. In various embodiments, the antibody is a polyclonal antibody or a monoclonal antibody. In one embodiment, the protein library is contacted *in vitro*. In one embodiment, the protein library is selected from the group consisting of a phage display library, a phagemid display library, and a ribosomal display library.

[0132] In one aspect, the invention provides a method of determining the rate of transfer of nucleic acid from a minicell to a cell, comprising (a) contacting the cell to the minicell, wherein the minicell comprises the nucleic acid, for a measured period of time; (b) separating minicells from the cells; (c) measuring the amount of nucleic acid in the cells, wherein the amount of nucleic acid in the cells over the set period of time is the rate of transfer of a nucleic acid from a minicell.

[0133] In one aspect, the invention provides a method of determining the amount of a nucleic acid transferred to a cell from a minicell, comprising (a) contacting the cell to the minicell, wherein the minicell comprises an expression element having eukaryotic expression sequences operably linked to an ORF encoding a detectable polypeptide, wherein the minicell displays a binding moiety, and wherein the binding moiety binds an epitope of the cell; and (b) detecting a signal from the detectable polypeptide, wherein

a change in the signal corresponds to an increase in the amount of a nucleic acid transferred to a cell.

[0134] In one embodiment, the cell is a eukaryotic cell. By way of non-limiting example, a eukaryotic cell can be a plant cell, a fungal cell, a unicellular eukaryote, an animal cell, a mammalian cell, a rat cell, a mouse cell, a primate cell or a human cell.

[0135] In one embodiment, the binding moiety is an antibody or antibody derivative. In one embodiment, the binding moiety is a single-chain antibody. In one embodiment, the binding moiety is an aptamer. In one embodiment, the binding moiety is an organic compound. In one embodiment, the detectable polypeptide is a fluorescent polypeptide.

[0136] In one aspect, the invention provides a method of detecting the expression of an expression element in a cell, comprising (a) contacting the cell to a minicell, wherein the minicell comprises an expression element having cellular expression sequences operably linked to an ORF encoding a detectable polypeptide, wherein the minicell displays a binding moiety, and wherein the binding moiety binds an epitope of the cell; (b) incubating the cell and the minicell for a period of time effective for transfer of nucleic acid from the minicell to the cell; and (c) detecting a signal from the detectable polypeptide, wherein an increase in the signal corresponds to an increase in the expression of the expression element.

[0137] In one embodiment, the cell is a eukaryotic cell and the expression sequences are eukaryotic expression sequences. In one embodiment, the eukaryotic cell is a mammalian cell. In one embodiment, the binding moiety is an antibody or antibody derivative. In one embodiment, the binding moiety is a single-chain antibody. In one embodiment, the binding moiety is an aptamer. In one embodiment, the binding moiety is an organic compound.

[0138] In a related aspect, the invention provides methods of detecting the transfer of a fusion protein from the cytosol to an organelle of a eukaryotic cell, comprising (a) contacting the cell to a minicell, wherein (i) the minicell comprises an expression element having eukaryotic expression sequences operably linked to an ORF encoding a fusion protein, wherein the fusion protein comprises a first polypeptide that comprises organellar delivery sequences, and a second polypeptide that comprises a detectable polypeptide; and (ii) the minicell displays a binding moiety that binds an epitope of the cell, or an epitope of an organelle; (b) incubating the cell and the minicell for a period of time effective for transfer of nucleic acid from the minicell to the cell and production of the fusion protein; and (c) detecting a signal from the detectable polypeptide, wherein a change in the signal corresponds to an increase in the amount of the fusion protein transferred to the organelle.

[0139] In one aspect, the invention provides a minicell comprising at least one nucleic acid, wherein the minicell displays a binding moiety directed to a target compound, wherein the binding moiety is selected from the group consisting of (i) a eukaryotic membrane protein; (ii) an archeobacterial membrane protein; (iii) an organellar membrane protein; and (iv) a fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one

membrane anchoring domain; and a second polypeptide, wherein the second polypeptide is not derived from a eubacterial protein and is neither a His tag nor a glutathione-S-transferase polypeptide, and wherein the polypeptide comprises a binding moiety.

[0140] In one embodiment, the nucleic acid comprises an expression construct comprising expression sequences operably linked to an ORF encoding a protein selected from the group consisting of (i) the eukaryotic membrane protein, (ii) the archeobacterial membrane protein, (iii) the organellar membrane protein; and (iv) the fusion protein.

[0141] In one embodiment, the nucleic acid comprises an expression construct comprising expression sequences operably linked to an ORF, wherein the ORF encodes a therapeutic polypeptide. In one embodiment, the therapeutic polypeptide is a membrane polypeptide. In one embodiment, the therapeutic polypeptide is a soluble polypeptide. In one embodiment, the soluble polypeptide comprises a cellular secretion sequence. In one embodiment, the expression sequences are inducible and/or repressible.

[0142] In one embodiment, the expression sequences are induced and/or derepressed when the binding moiety displayed by the minicell binds to its target compound. In one embodiment, the nucleic acid comprises an expression construct comprising expression sequences operably linked to an ORF, wherein the ORF encodes a polypeptide having an amino acid sequence that facilitates cellular transfer of a biologically active compound contained within or displayed by the minicell. In one embodiment, the membrane of the minicell comprises a system for transferring a molecule from the interior of a minicell into the cytoplasm of the cell. In one embodiment, the system for transferring a molecule from the interior of a minicell into the cytoplasm of the cell is a Type III secretion system.

[0143] In one aspect, the invention provides a method of introducing a nucleic acid into a cell, comprising contacting the cell with a minicell that comprises the nucleic acid, wherein the minicell displays a binding moiety, wherein the binding moiety is selected from the group consisting of (i) a eukaryotic membrane protein; (ii) an archeobacterial membrane protein; (iii) an organellar membrane protein; and (iv) a fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and a second polypeptide, wherein the second polypeptide is not derived from a eubacterial protein and is neither a His tag nor a glutathione-S-transferase polypeptide, and wherein the polypeptide comprises a binding moiety; and wherein the binding moiety binds an epitope of the cell.

[0144] In one embodiment, the nucleic acid comprises an expression construct comprising expression sequences operably linked to an ORF encoding a protein selected from the group consisting of (i) the eukaryotic membrane protein, (ii) the archeobacterial membrane protein, (iii) the organellar membrane protein; and (iv) a fusion protein.

[0145] In one embodiment, the nucleic acid comprises an expression construct comprising expression sequences operably linked to an ORF, wherein the ORF encodes a therapeutic polypeptide. In one embodiment, the expression sequences are inducible and/or derepressible. In one

embodiment, the expression sequences are induced or derepressed when the binding moiety displayed by the minicell binds its target compound. In one embodiment, the expression sequences are induced or derepressed by a transactivation or transrepression event. In one embodiment, the nucleic acid comprises an expression construct comprising expression sequences operably linked to an ORF, wherein the ORF encodes a polypeptide having an amino acid sequence that facilitates cellular transfer of a biologically active compound contained within or displayed by the minicell.

**[0146]** In one aspect, the invention provides a minicell comprising a nucleic acid, wherein the nucleic acid comprises eukaryotic expression sequences and eubacterial expression sequences, each of which is independently operably linked to an ORF.

**[0147]** In one embodiment, the minicell displays a binding moiety. In one embodiment, the eubacterial expression sequences are induced and/or derepressed when the binding moiety is in contact with a target cell. In one embodiment, the eukaryotic expression sequences are induced and/or derepressed when the nucleic acid is in the cytoplasm of a eukaryotic cell. In one embodiment, the protein encoded by the ORF comprises eubacterial or eukaryotic secretion sequences.

**[0148]** In one aspect, the invention provides a minicell comprising a first and second nucleic acid, wherein the first nucleic acid comprises eukaryotic expression sequences operably linked to a first ORF, and a second nucleic acid, wherein the second nucleic acid comprises eubacterial expression sequences operably linked to a second ORF.

**[0149]** In one embodiment, the minicell displays a binding moiety. In one embodiment, the eubacterial expression sequences are induced and/or derepressed when the binding moiety is in contact with a target cell. In one embodiment, the eukaryotic expression sequences are induced and/or derepressed when the nucleic acid is in the cytoplasm of a eukaryotic cell. In one embodiment, the protein encoded by the first ORF comprises eukaryotic secretion sequences and/or the protein encoded by the second ORF comprises eubacterial secretion sequences.

**[0150]** In one aspect, the invention provides a method of introducing into and expressing a nucleic acid in an organism, comprising contacting a minicell to a cell of the organism, wherein the minicell comprises the nucleic acid.

**[0151]** In one embodiment, the minicell displays a binding moiety. In one embodiment, the nucleic acid comprises a eukaryotic expression construct, wherein the eukaryotic expression construct comprises eukaryotic expression sequences operably linked to an ORF. In one embodiment, the ORF encodes a protein selected from the group consisting of a membrane protein, a soluble protein and a protein comprising eukaryotic secretion signal sequences. In one embodiment, the nucleic acid comprises a eubacterial expression construct, wherein the eubacterial expression construct comprises eubacterial expression sequences operably linked to an ORF. In one embodiment, the minicell displays a binding moiety, wherein the eubacterial expression sequences are induced and/or derepressed when the binding moiety is in contact with a target cell. In one embodiment, the protein encoded by the ORF comprises

eubacterial secretion sequences. In one aspect, the invention provides a minicell comprising a crystal of a membrane protein. In one embodiment, the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast. In one embodiment, the membrane protein is a receptor. In one embodiment, the receptor is a G-protein coupled receptor. In one embodiment, the crystal is displayed.

**[0152]** In a related aspect, the invention provides a minicell membrane preparation comprising a crystal of a membrane protein.

**[0153]** In one embodiment, the membrane protein is a fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain, and a second polypeptide. In one embodiment, the crystal is a crystal of the second polypeptide. In one embodiment, the crystal is displayed.

**[0154]** In one aspect, the invention provides a method of determining the three-dimensional structure of a membrane protein, comprising preparing a crystal of the membrane protein in a minicell, and determining the three-dimensional structure of the crystal.

**[0155]** In one aspect, the invention provides a method for identifying ligand-interacting atoms in a defined three-dimensional structure of a target protein, comprising (a) preparing one or more variant proteins of a target protein having a known or predicted three-dimensional structure, wherein the target protein binds a preselected ligand; (b) expressing and displaying a variant protein in a minicell; and (c) determining if a minicell displaying the variant protein binds the preselected ligand with increased or decreased affinity as compared to the binding of the preselected ligand to the target protein.

**[0156]** In one embodiment, the ligand is a protein that forms a multimer with the target protein, and the ligand interacting atoms are atoms in the defined three-dimensional structure are atoms that are involved in protein-protein interactions. In one embodiment, the ligand is a compound that induces a conformational change in the target protein, and the defined three-dimensional structure is the site of the conformational change. In one embodiment, the method for identifying ligands of a target protein, further comprising identifying the chemical differences in the variant proteins as compared to the target protein. In one embodiment, the invention further comprises mapping the chemical differences onto the defined three-dimensional structure, and correlating the effect of the chemical differences on the defined three-dimensional structure. In one embodiment, the target protein is a wild-type protein. In one aspect, the invention provides a minicell library, comprising two or more minicells, wherein each minicell comprises a different exogenous protein. In one embodiment, the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast. In one embodiment, the exogenous protein is a displayed protein. In one embodiment, the exogenous protein is a membrane protein. In one embodiment, the membrane protein is a receptor. In one embodiment, the protein is a soluble protein that is contained within or secreted from the minicell. In one embodiment, minicells within the library comprise an expression element that comprises expression sequences operably linked to a nucleic acid having an ORF that encodes the exogenous protein. In one embodiment, the

nucleic acid has been mutagenized; the mutagenesis can be site-directed or random. In one embodiment, an active site of the exogenous protein has a known or predicted three-dimensional structure, and the a portion of the ORF encoding the active site has been mutagenized. In one embodiment, each of the minicells comprises an exogenous protein that is a variant of a protein having a known or predicted three-dimensional structure.

[0157] In one aspect, the invention provides a minicell library, comprising two or more minicells, wherein each minicell comprises a different fusion protein, each of the fusion protein comprising a first polypeptide that is a constant polypeptide, wherein the constant polypeptide comprises at least one transmembrane domain or at least one membrane anchoring domain, and a second polypeptide, wherein the second polypeptide is a variable amino acid sequence that is different in each fusion proteins. In one embodiment, minicells within the library comprise an expression element that comprises expression sequences operably linked to a nucleic acid having an ORF that encodes the fusion protein. In one embodiment, the second polypeptide of the fusion protein is encoded by a nucleic acid that has been cloned. In one embodiment, each of the second polypeptide of each of the fusion proteins comprises a variant of an amino acid sequence from a protein having a known or predicted three-dimensional structure.

[0158] In one aspect, the invention provides a minicell library, comprising two or more minicells, wherein each minicell comprises a constant protein that is present in each minicell and a variable protein that differs from minicell to minicell. In one embodiment, one of the constant and variable proteins is a receptor, and the other of the constant and variable proteins is a co-receptor. In one embodiment, each of the constant and variable proteins is different from each other and is a factor in a signal transduction pathway. In one embodiment, one of the constant and variable proteins is a G-protein, and the other of the constant and variable proteins is a G-protein coupled receptor.

[0159] In one embodiment, one of the constant and variable proteins comprises a first transrepression domain, and the other of the constant and variable comprises a second transrepression domain, wherein the transrepression domains limit or block expression of a reporter gene when the constant and variable proteins associate with each other.

[0160] In one embodiment, one of the constant and variable proteins comprises a first transactivation domain, and the other of the constant and variable comprises a second transactivation domain, wherein the transactivation domains stimulate expression of a reporter gene when the constant and variable proteins associate with each other.

[0161] In one aspect, the invention provides a method of identifying a nucleic acid that encodes a protein that binds to or chemically alters a preselected ligand, comprising (a) separately contacting the ligand with individual members of a minicell library, wherein minicells in the library comprise expression elements, wherein the expression elements comprise DNA inserts, wherein an ORF in the DNA insert is operably linked to expression sequences, in order to generate a series of reaction mixes, each reaction mix comprising a different member of the minicell library; (b) incubating the reaction mixes, thereby allowing a protein that binds to or chemically alters the preselected ligand to bind or chemi-

cally alter the preselected ligand; (c) detecting a change in a signal from reaction mixes in which the ligand has been bound or chemically altered; (d) preparing DNA from reaction mixes in which the ligand has been bound or chemically altered; wherein the DNA is a nucleic acid that encodes a protein that binds to or chemically alters the preselected ligand.

[0162] In one embodiment, the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast. In one embodiment, the preselected ligand is a biologically active compound. In one embodiment, the preselected ligand is a therapeutic drug. In one embodiment, a protein that binds or chemically alters the preselected ligand is a target protein for compounds that are therapeutic for a disease that is treated by administering the drug to an organism in need thereof. In one embodiment, the preselected ligand is detectably labeled, the minicell comprises a detectable compound, and/or a chemically altered derivative of the protein is detectably labeled.

[0163] In one aspect, the invention provides a method of determining the amino acid sequence of a protein that binds or chemically alters a preselected ligand, comprising: (a) contacting the ligand with a minicell library, wherein minicells in the library comprise expression elements, wherein the expression elements comprise DNA inserts, wherein an ORF in the DNA insert is operably linked to expression sequences; (b) incubating the mixture of ligand and minicells, under conditions which allow complexes comprising ligands and minicells to form and/or chemical reactions to occur; (c) isolating or identifying the complexes from the ligand and the mixture of ligand and minicells; (d) preparing DNA from an expression element found in one or more of the complexes, or in a minicell thereof; (e) determining the nucleotide sequence of the ORF in the DNA; and (f) generating an amino sequence by in silico translation, wherein the amino acid sequence is or is derived from a protein that binds or chemically alters a preselected ligand.

[0164] In one embodiment, the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast. In one embodiment, the DNA is prepared by isolating DNA from the complexes, or in a minicell thereof. In one embodiment, the DNA is prepared by amplifying DNA from the complexes, or in a minicell thereof. In one embodiment, the protein is a fusion protein. In one embodiment, the protein is a membrane or a soluble protein. In one embodiment, the protein comprises secretion sequences. In one embodiment, the preselected ligand is a biologically active compound. In one embodiment, the preselected ligand is a therapeutic drug. In one embodiment, the preselected ligand is a therapeutic drug, and the protein that binds the preselected ligand is a target protein for compounds that are therapeutic for a disease that is treated by administering the drug to an organism in need thereof.

[0165] In one aspect, the invention provides a method of identifying a nucleic acid that encodes a protein that inhibits or blocks an agent from binding to or chemically altering a preselected ligand, comprising: (a) separately contacting the ligand with individual members of a minicell library, wherein minicells in the library comprise expression elements, wherein the expression elements comprise DNA inserts, wherein an ORF in the DNA insert is operably linked to expression sequences, in order to generate a series of

reaction mixes, each reaction mix comprising a different member of the minicell library; (b) incubating the reaction mixes, thereby allowing a protein that binds to or chemically alters the preselected ligand to bind or chemically alter the preselected ligand; (c) detecting a change in a signal from reaction mixes in which the ligand has been bound or chemically altered; (d) preparing DNA from reaction mixes in which the change in signal ligand has been bound or chemically altered; wherein the DNA is a nucleic acid that encodes a protein that inhibits or blocks the agent from binding to or chemically altering the preselected ligand. In one embodiment, the minicell is a eubacterial minicell, a poroplast, a spheroplast or a protoplast. In one embodiment, the DNA has a nucleotide sequence that encodes the amino acid sequence of the protein that inhibits or blocks the agent from binding to or chemically altering the preselected ligand. In one embodiment, a protein that binds or chemically alters the preselected ligand is a target protein for compounds that are therapeutic for a disease that is treated by administering the drug to an organism in need thereof.

**[0166]** In one aspect, the invention provides a method of identifying an agent that effects the activity of a protein, comprising contacting a library of two or more candidate agents with a minicell comprising the protein or a polypeptide derived from the protein, assaying the effect of candidate agents on the activity of the protein, and identifying agents that effect the activity of the protein.

**[0167]** In one embodiment, the protein or the polypeptide derived from the protein is displayed on the surface of the minicell. In one embodiment, the protein is a membrane protein. In one embodiment, the membrane protein is selected from the group consisting of a receptor, a channel protein and an enzyme. In one embodiment, the activity of a protein is a binding activity or an enzymatic activity. In one embodiment, the library of compounds is a protein library. In one embodiment, the protein library is selected from the group consisting of a phage display library, a phagemid display library, and a ribosomal display library. In one embodiment, the library of compounds is a library of aptamers. In one embodiment, the library of compounds is a library of small molecules.

**[0168]** In one aspect, the invention provides a method of identifying an agent that effects the activity of a protein domain containing a library of two or more candidate agents with a minicell displaying a membrane fusion protein, the fusion protein comprising a first polypeptide, the first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain, and a second polypeptide, wherein the second polypeptide comprises the protein domain.

**[0169]** In one aspect, the invention provides a method of identifying undesirable side-effects of a biologically active compound that occur as a result of binding of the compound to a protein, wherein binding a compound to the protein is known to result in undesirable side effects, comprising contacting a minicell that comprises the protein to the biologically active compound. In one embodiment, the invention provides comprises characterizing the binding of the biologically active compound to the protein. In one embodiment, the invention provides comprises characterizing the effect of the biologically active compound on the activity of the protein.

**[0170]** In one aspect, the invention provides a method for identifying an agent that effects the interaction of a first signaling protein with a second signaling protein, comprising (a) contacting a library of compounds with a minicell, wherein the minicell comprises: (i) a first protein comprising the first signaling protein and a first trans-acting regulatory domain; (ii) a second protein comprising the second signaling protein and a second trans-acting regulatory domain; and (iii) a reporter gene, the expression of which is modulated by the interaction between the first trans-acting regulatory domain and the second trans-acting regulatory domain; and (b) detecting the gene product of the reporter gene.

**[0171]** In one embodiment, the trans-acting regulatory domains are transactivation domains. In one embodiment, the trans-acting regulatory domains are transrepression domains.

**[0172]** In one embodiment, the reporter gene is induced by the interaction of the first trans-acting regulatory domain and the second trans-acting regulatory domain. In one embodiment, the agent that effects the interaction of the first signaling protein with the second signaling protein is an agent that causes or promotes the interaction. In one embodiment, the reporter gene is repressed by the interaction of the first trans-acting regulatory domain and the second trans-acting regulatory domain. In one embodiment, the agent that effects the interaction of the first signaling protein with the second signaling protein is an agent that inhibits or blocks the interaction.

**[0173]** In one embodiment, the first signaling protein is a GPCR. In one embodiment, the GPCR is an Edg receptor or a ScAMPER.

**[0174]** In one embodiment, the second signalling protein is a G-protein. In related embodiments, G-protein is selected from the group consisting of G-alpha-i, G-alpha-s, G-alpha-q, G-alpha-12/13 and Go. In one embodiment, the library of compounds is a protein library. In one embodiment, the protein library is selected from the group consisting of a phage display library, a phagemid display library, and a ribosomal display library. In one embodiment, the library of compounds is a library of aptamers. In one embodiment, the library of compounds is a library of small molecules.

**[0175]** In one aspect, the invention provides a method for identifying an agent that effects the interaction of a first signaling protein with a second signaling protein, comprising contacting a library of two or more candidate agents with a minicell, wherein the minicell comprises (a) a first fusion protein comprising the first signaling protein and a first detectable domain; and (b) a second fusion protein comprising the second signaling protein and a second detectable domain, wherein a signal is generated when the first and second signaling proteins are in close proximity to each other, and detecting the signal.

**[0176]** In one embodiment, the signal is fluorescence. In one embodiment, the first detectable domain and the second detectable domain are fluorescent and the signal is generated by FRET. In one embodiment, the first and second detectable domains are independently selected from the group consisting of a green fluorescent protein, a blue-shifted green fluorescent protein, a cyan-shifted green fluorescent protein; a red-shifted green fluorescent protein; a yellow-shifted green fluorescent protein, and a red fluorescent protein,

wherein the first fluorescent domain and the second fluorescent domain are not identical.

[0177] In one aspect, the invention provides a method of bioremediation, the method comprising contacting a composition that comprises an undesirable substance with a minicell, wherein the minicell alters the chemical structure and/or binds the undesirable substance.

[0178] In one aspect, the invention provides a method of bioremediation, the method comprising contacting a composition that comprises an undesirable substance with a minicell, wherein the minicell comprises an agent that alters the chemical structure of the undesirable substance. In one embodiment, the agent that alters the chemical structure of the undesirable substance is an inorganic catalyst. In one embodiment, the agent that alters the chemical structure of the undesirable substance is an enzyme. In one embodiment, the enzyme is a soluble protein contained within the minicell. In one embodiment, the enzyme is a secreted protein. In one embodiment, the enzyme is a membrane protein. In one embodiment, the membrane enzyme is selected from the group consisting of a cytochrome P450, an oxidoreductase, a transferase, a hydrolase, a lyase, an isomerase, a ligase and a synthetase. In one embodiment, the agent that alters the chemical structure of the undesirable substance is a fusion protein comprising a first polypeptide that comprises a transmembrane domain or at least one membrane-anchoring domain, and a second polypeptide, wherein the second polypeptide is an enzyme moiety.

[0179] In one aspect, the invention provides a method of bioremediation, the method comprising contacting a composition that comprises an undesirable substance with a minicell, wherein the minicell comprises an agent that binds an undesirable substance. In one embodiment, the undesirable substance binds to and is internalized by the minicell or is otherwise inactivated by selective absorption. In one embodiment, the agent that binds the undesirable substance is a secreted soluble protein. In one embodiment, the secreted protein is a transport accessory protein. In one embodiment, the agent that binds the undesirable substance is a membrane protein. In one embodiment, the undesirable substance is selected from the group consisting of a toxin, a pollutant and a pathogen. In one embodiment, the agent that binds the undesirable substance is a fusion protein comprising a first polypeptide that comprises a transmembrane domain or at least one membrane-anchoring domain, and a second polypeptide, wherein the second polypeptide is a binding moiety. In one embodiment, wherein the binding moiety is selected from the group consisting of an antibody, an antibody derivative, the active site of a non-enzymatically active mutant enzyme, a single-chain antibody and an aptamer.

[0180] In one aspect, the invention provides a minicell-producing parent cell, wherein the parent cell comprises one or more of the following (a) an expression element that comprises a gene operably linked to expression sequences that are inducible and/or repressible, wherein induction or repression of the gene regulates the copy number of an episomal expression construct; (b) a mutation in an endogenous gene, wherein the mutation regulates the copy number of an episomal expression construct; (c) an expression element that comprises a gene operably linked to expression sequences that are inducible and/or repressible, wherein

induction or repression of the gene causes or enhances the production of minicells; and (d) a mutation in an endogenous gene, wherein the mutation causes or enhances minicell production.

[0181] In one embodiment, the invention comprises an episomal expression construct. In one embodiment, the invention further comprises a chromosomal expression construct. In one embodiment, the expression sequences of the expression construct are inducible and/or repressible. In one embodiment, the minicell-producing parent cell comprises a biologically active compound. In one embodiment, the gene that causes or enhances the production of minicells has a gene product that is involved in or regulates DNA replication, cellular division, cellular partitioning, septation, transcription, translation, or protein folding.

[0182] In one aspect, the invention provides a minicell-producing parent cell, wherein the parent cell comprises an expression construct, wherein the expression construct comprises expression sequences operably linked to an ORF that encodes a protein, and a regulatory expression element, wherein the regulatory expression element comprises expression sequences operably linked to a regulatory gene that encodes a factor that regulates the expression of the ORF. In one embodiment, the expression sequences of the expression construct are inducible and/or repressible. In one embodiment, the expression sequences of the regulatory expression construct are inducible and/or repressible. In one embodiment, one or more of the expression element or the regulatory expression element is located on a chromosome of the parent cell. In one embodiment, one or more of the expression element or the regulatory expression element is located on an episomal expression construct. In one embodiment, both of the expression element and the regulatory expression element are located on an episomal expression construct, and one or both of the expression element and the regulatory expression element segregates into minicells produced from the parent cell. In one embodiment, the minicell-producing parent cell comprises a biologically active compound. In one embodiment, the biologically active compound segregates into minicells produced from the parent cell. In one embodiment, the ORF encodes a membrane protein or a soluble protein. In one embodiment, the protein comprises secretion sequences. In one embodiment, the gene product of the gene regulates the expression of the ORF. In one embodiment, the gene product is a transcription factor. In one embodiment, the gene product is a RNA polymerase. In one embodiment, the parent cell is MC-T7.

[0183] In one aspect, the invention provides a minicell comprising a biologically active compound, wherein the minicell displays a binding moiety, wherein the minicell selectively absorbs and/or internalizes an undesirable compound, and the minicell is a poroplast, spheroplast or protoplast. In one embodiment, the binding moiety is selected from the group consisting of an antibody, an antibody derivative, a receptor and an active site of a non-catalytic derivative of an enzyme. In one embodiment, the binding moiety is a single-chain antibody. In one embodiment, the binding moiety is directed to a ligand selected from the group consisting of an epitope displayed on a pathogen, an epitope displayed on an infected cell and an epitope displayed on a hyperproliferative cell. In one embodiment, the biologically active compound is selected from the group consisting of a radioisotope, a polypeptide, a nucleic acid



and a small molecule. In one embodiment, a ligand binds to and is internalized by the minicell or is otherwise inactivated by selective absorption. In one embodiment, the invention provides a pharmaceutical composition comprising the minicell. In one aspect, the invention provides a method of reducing the free concentration of a substance in a composition, wherein the substance displays a ligand specifically recognized by a binding moiety, comprising contacting the composition with a minicell that displays the binding moiety, wherein the binding moiety binds the substance, thereby reducing the free concentration of the substance in the composition. In one embodiment, the substance is selected from the group consisting of a nucleic acid, a lipid, a polypeptide, a radioactive compound, an ion and a small molecule. In one embodiment, the binding moiety is selected from the group consisting of an antibody, an antibody derivative, a channel protein and a receptor.

[0184] In one embodiment, the composition is present in an environment including but not limited to water, air or soil. In one embodiment, the composition is a biological sample from an organism, including but not limited to blood, serum, plasma, urine, saliva, a biopsy sample, feces, tissue and a skin patch. In one embodiment, the substance binds to and is internalized by the minicell or is otherwise inactivated by selective absorption. In one embodiment, the biological sample is returned to the organism after being contacting to the minicell.

[0185] For a better understanding of the present invention, reference is made to the accompanying detailed description and its scope will be pointed out in the appended claims. All references cited herein are hereby incorporated by reference.

#### BRIEF DESCRIPTION OF THE DRAWINGS

[0186] FIG. 1 is a Western blot in which Edg-1-6× His and Edg-3-6× His proteins expressed in minicells produced from MC-T7 cells.

[0187] FIG. 2 shows induction of MalE(L)-NTR in isolated minicells.

#### ABBREVIATIONS AND DEFINITIONS

[0188] For brevity's sake, the single-letter amino acid abbreviations are used in some instances herein. Table 1 describes the correspondence between the 1- and 3-letter amino acid abbreviations.

TABLE 1

THREE- AND ONE- LETTER ABBREVIATIONS FOR AMINO ACIDS		
Amino acid	Three-letter abbreviation	One-letter symbol
Alanine	Ala	A
Arginine	Arg	R
Asparagine	Asn	N
Aspartic Acid	Asp	D
Cysteine	Cys	C
Glutamine	Gln	Q
Glutamic acid	Glu	E
Glycine	Gly	G
Histidine	His	H
Isoleucine	Ile	I
Leucine	Leu	L
Lysine	Lys	K
Methionine	Met	M

TABLE 1-continued

THREE- AND ONE- LETTER ABBREVIATIONS FOR AMINO ACIDS		
Amino acid	Three-letter abbreviation	One-letter symbol
Phenylalanine	Phe	F
Proline	Pro	P
Serine	Ser	S
Threonine	Thr	T
Tryptophan	Trp	W
Tyrosine	Tyr	Y
Valine	Val	V

[0189] A “conjugatable compound” or “attachable compound” is capable of being attached to another compound. The terms “conjugated to” and “cross-linked with” indicate that the conjugatable compound is in the state of being attached to another compound. A “conjugate” is the compound formed by the attachment of a conjugatable compound or conjugatable moiety to another compound.

[0190] “Culturing” signifies incubating a cell or organism under conditions wherein the cell or organism can carry out some, if not all, biological processes. For example, a cell that is cultured may be growing or reproducing, or it may be non-viable but still capable of carrying out biological and/or biochemical processes such as replication, transcription, translation, etc.

[0191] An agent is said to have been “purified” if its concentration is increased, and/or the concentration of one or more undesirable contaminants is decreased, in a composition relative to the composition from which the agent has been purified. Purification thus encompasses enrichment of an agent in a composition and/or isolation of an agent therefrom.

[0192] A “solid support” is any solid or semisolid composition to which an agent can be attached or contained within. Common forms of solid support include, but are not limited to, plates, tubes, and beads, all of which could be made of glass or another suitable material, e.g., polystyrene, nylon, cellulose acetate, nitrocellulose, and other polymers. Semisolids and gels that minicells are suspended within are also considered to be solid supports. A solid support can be in the form of a dipstick, flow-through device, or other suitable configuration.

[0193] A “mutation” is a change in the nucleotide sequence of a gene relative to the sequence of the “wild-type” gene. Reference wild-type eubacterial strains are those that have been cultured in vitro by scientists for decades; for example, a wild-type strain of *Escherichia coli* is *E. coli* K-12. Mutations include, but are not limited to, point mutations, deletions, insertions and translocations.

[0194] A “trans-acting regulatory domain” is a regulatory part of a protein that is expressed from a gene that is not adjacent to the site of regulatory effect. Trans-acting domains can activate or stimulate (transactivate), or limit or block (transrepress) the gene in question.

[0195] A “reporter gene” refers to a gene that is operably linked to expression sequences, and which expresses a gene product, typically a detectable polypeptide, the production and detection of which is used as a measure of the robustness and/or control of expression.

[0196] A “detectable compound” or “detectable moiety” produces a signal that can be detected by spectroscopic, photochemical, biochemical, immunochemical, electromagnetic, radiochemical, or chemical means such as fluorescence, chemifluorescence, or chemiluminescence, or any other appropriate means. A “radioactive compound” or “radioactive composition” has more than the natural (environmental) amount of one or more radioisotopes.

[0197] By “displayed” it is meant that a portion of the membrane protein is present on the surface of a cell or minicell, and is thus in contact with the external environment of the cell or minicell. The external, displayed portion of a membrane protein is an “extracellular domain” or a “displayed domain.” A membrane protein may have more than one displayed domain, and a minicell of the invention may display more than one membrane protein.

[0198] A “domain” or “protein domain” is a region of a molecule or structure that shares common physical and/or chemical features. Non-limiting examples of protein domains include hydrophobic transmembrane or peripheral membrane binding regions, globular enzymatic or receptor regions, and/or nucleic acid binding domains.

[0199] A “transmembrane domain” spans a membrane, a “membrane anchoring domain” is positioned within, but does not traverse, a membrane. An “extracellular” or “displayed” domain is present on the exterior of a cell, or minicell, and is thus in contact with the external environment of the cell or minicell.

[0200] A “eukaryote” is as the term is used in the art. A eukaryote may, by way of non-limiting example, be a fungus, a unicellular eukaryote, a plant or an animal. An animal may be a mammal, such as a rat, a mouse, a rabbit, a dog, a cat, a horse, a cow, a pig, a simian or a human.

[0201] A “eukaryotic membrane” is a membrane found in a eukaryote. A eukaryotic membrane may, by way of non-limiting example, be a cytoplasmic membrane, a nuclear membrane, a nucleolar membrane, a membrane of the endoplasmic reticulum (ER), a membrane of a Golgi body, a membrane of a lysosome, a membrane of a peroxisome, a caveolar membrane, or an inner or outer membrane of a mitochondrion, chloroplast or plastid.

[0202] The term “endogenous” refers to something that is normally found in a cell as that cell exists in nature.

[0203] The term “exogenous” refers to something that is not normally found in a cell as that cell exists in nature.

[0204] A “gene” comprises (a) nucleotide sequences that either (i) act as a template for a nucleic acid gene product, or (ii) that encode one or more open reading frames (ORFs); and (b) expression sequences operably linked to (1) or (2). When a gene comprises an ORF, it is a “structural gene.”

[0205] By “immunogenic,” it is meant that a compound elicits production of antibodies or antibody derivatives and, additionally or alternatively, a T-cell mediated response, directed to the compound or a portion thereof. The compound is an “immunogen.”

[0206] A “ligand” is a compound, composition or moiety that is capable of specifically bound by a binding moiety, including without limitation, a receptor and an antibody or antibody derivative.

[0207] A “membrane protein” is a protein found in whole or in part in a membrane. Typically, a membrane protein has (1) at least one membrane anchoring domain, (2) at least one transmembrane domain, or (3) at least one domain that interacts with a protein having (1) or (2).

[0208] An “ORF” or “open reading frame” is a nucleotide sequence that encodes an amino acid sequence of a known, predicted or hypothetical polypeptide. An ORF is bounded on its 5' end by a start codon (usually ATG) and on its 3' end by a stop codon (i.e., TAA or TGA). An ORF encoding a 10 amino acid sequence comprises 33 nucleotides (3 for each of 10 amino acids and 3 for a stop codon). ORFs can encode amino acid sequences that comprise from 10, 25, 50, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 600, 700, 800, 900 or more amino acids. The terms “Eubacteria” and “prokaryote” are used herein as these terms are used by those in the art. The terms “eubacterial” and “prokaryotic” encompasses Eubacteria, including both gram-negative and gram-positive bacteria, prokaryotic viruses (e.g., bacteriophage), and obligate intracellular parasites (e.g., Rickettsia, Chlamydia, etc.).

[0209] An “active site” is any portion or region of a molecule required for, or that regulates, an activity of the molecule. In the case of a protein, an active site can be a binding site for a ligand or a substrate, an active site of enzyme, a site that directs or undergoes conformational change in response to a signal, or a site of post-translational modification of a protein.

[0210] In a poroplast, the eubacterial outer membrane (OM) and LPS have been removed. In a spheroplast, portions of a disrupted eubacterial OM and/or disrupted cell wall either may remain associated with the inner membrane of the minicell, but the membrane is nonetheless porous because the permeability of the disrupted OM has been increased. A membrane is to be “disrupted” when the membrane’s structure has been treated with an agent, or incubated under conditions, that leads to the partial degradation of the membrane, thereby increasing the permeability thereof. In contrast, a membrane that has been “degraded” is essentially, for the applicable intents and purposes, removed. In preferred embodiments, irrespective of the condition of the OM and cell wall, the eubacterial inner membrane is not disrupted, and membrane proteins displayed on the inner membrane are accessible to compounds that are brought into contact with the minicell, poroplast, spheroplast, protoplast or cellular poroplast, as the case may be.

[0211] Host cells (and/or minicells) harboring an expression construct are components of expression systems.

[0212] An “expression vector” is an artificial nucleic acid molecule into which an exogenous ORF encoding a protein, or a template of a bioactive nucleic acid can be inserted in such a manner so as to be operably linked to appropriate expression sequences that direct the expression of the exogenous gene. Preferred expression vectors are episomal vectors that can replicate independently of chromosomal replication.

[0213] By the term “operably linked” it is meant that the gene products encoded by the non-vector nucleic acid sequences are produced from an expression element in vivo.

[0214] The term “gene product” refers to either a nucleic acid (the product of transcription, reverse transcription, or

replication) or a polypeptide (the product of translation) that is produced using the non-vector nucleic acid sequences as a template.

[0215] An “expression construct” is an expression vector into which a nucleotide sequence of interest has been inserted in a manner so as to be positioned to be operably linked to the expression sequences present in the expression vector. Preferred expression constructs are episomal.

[0216] An “expression element” is a nucleic acid having nucleotide sequences that are present in an expression construct but not its cognate expression vector. That is, an expression element for a polypeptide is a nucleic acid that comprises an ORF operably linked to appropriate expression sequences. An expression element can be removed from its expression construct and placed in other expression vectors or into chromosomal DNA.

[0217] “Expression sequences” are nucleic acid sequences that bind factors necessary for the expression of genes that have been inserted into an expression vector. An example of an expression sequence is a promoter, a sequence that binds RNA polymerase, which is the enzyme that produces RNA molecules using DNA as a template. An example of an expression sequence that is both inducible and repressible is L-arabinose operon (araC). See Schleif R. Regulation of the L-arabinose operon of *Escherichia coli*. Trends Genet. 2000 December;16(12):559-65.

[0218] In the present disclosure, “a nucleic acid” or “the nucleic acid” refers to a specific nucleic acid molecule. In contrast, the term “nucleic acid” refers to any collection of diverse nucleic acid molecules, and thus signifies that any number of different types of nucleic acids are present. By way of non-limiting example, a nucleic acid may be a DNA, a dsRNA, a tRNA (including a rare codon usage tRNA), an mRNA, a ribosomal RNA (rRNA), a peptide nucleic acid (PNA), a DNA:RNA hybrid, an antisense oligonucleotide, a ribozyme, or an aptamer.

#### DETAILED DESCRIPTION OF THE INVENTION

[0219] The invention described herein is drawn to compositions and methods for the production of achromosomal archaeobacterial, eubacterial and anucleate eukaryotic cells that are used for diagnostic and therapeutic applications, for drug discovery, and as research tools.

[0220] The general advantage of minicells over cell-based expression systems (e.g., eucaryotic cells or bacterial expression systems) is that one may express heterologous membrane bound proteins or over express endogenous membrane bound proteins, cytoplasmic or secreted soluble proteins, or small molecules on the cytoplasmic or extracellular surfaces of the minicells that would otherwise be toxic to live cells. Minicells are also advantageous for proteins that require a particular lipid environment for proper functioning because it is very manipulatable in nature. Other advantages include the stability of the minicells due to the lack of toxicity, the high level of expression that can be achieved in the minicell, and the efficient flexible nature of the minicell expression system. Such minicells could be used for in vivo targeting or for selective absorption (i.e., molecular “sponges”) and that these molecules can be expressed and “displayed” at high levels. Minicells can also

be used to display proteins for low, medium, high, and ultra high throughput screening, crystal formation for structure, determination, and for in vitro research use only applications such as transfection. Minicells expressing proteins or small molecules, radioisotopes, image-enhancing reagents can be used for in vivo diagnostics and for in vitro diagnostic and assay platforms. Also, soluble and/or membrane associated signaling cascade elements may be reconstituted in minicells producing encapsulated devices to follow extracellular stimulation events using cytoplasmic reporter events, e.g. transactivation resulting from dimerization of dimerization dependant transcriptional activation or repression of said reporter.

[0221] Regarding protein expression, minicells can be engineered to express one or more recombinant proteins in order to produce more protein per surface area of the particle (at least 10x more protein per unit surface area of protein). The proteins or small molecules that are “displayed” on the minicell surfaces can have therapeutic, discovery or diagnostic benefit either when injected into a patient or used in a selective absorption mode during dialysis. In vitro assays include drug screening and discovery, structural proteomics, and other functional proteomics applications. Proteins that are normally soluble can be tethered to membrane anchoring domains or membrane proteins can be expressed for the purpose of displaying these proteins on the surfaces of the minicell particle in therapeutic, discovery, and diagnostic modes. The types of proteins that can be displayed include but are not limited to receptors (e.g., GPCRs, sphingolipid receptors, neurotransmitter receptors, sensory receptors, growth factor receptors, hormone receptors, chemokine receptors, cytokine receptors, immunological receptors, and complement receptors, FC receptors), channels (e.g., potassium channels, sodium channels, calcium channels.), pores (e.g., nuclear pore proteins, water channels), ion and other pumps (e.g., calcium pumps, proton pumps), exchangers (e.g., sodium/potassium exchangers, sodium/hydrogen exchangers, potassium/hydrogen exchangers), electron transport proteins (e.g., cytochrome oxidase), enzymes and kinases (e.g., protein kinases, ATPases, GTPases, phosphatases, proteases.), structural/linker proteins (e.g., Caveolins, clathrin), adapter proteins (e.g., TRAD, TRAP, FAN), chemotactic/adhesion proteins (e.g., ICAM11, selectins, CD34, VCAM-1, LFA-1, VLA-1), and chimeric/fusion proteins (e.g., proteins in which a normally soluble protein is attached to a transmembrane region of another protein). As a non-limiting example, the small molecules that can be tethered and displayed on the surfaces of the minicells can be carbohydrates (e.g., monosaccharides), bioactive lipids (e.g., lysosphingolipids, PAF, lysophospholipids), drugs (e.g., antibiotics, ion channel activators/inhibitors, ligands for receptors and/or enzymes), nucleic acids (e.g., synthetic oligonucleotides), fluorophores, metals, or inorganic and organic small molecules typically found in combinatorial chemistry libraries. Minicells may either contain (encapsulate) or display on their surfaces radionuclides or image-enhancing reagents both of which could be used for therapeutic and/or diagnostic benefit in vivo or for in vitro assays and diagnostic platforms.

[0222] For in vivo therapeutic uses, minicells can express proteins and/or display small molecules on their surfaces that would either promote an immune response and passage through the RES system (e.g., to eliminate the minicell and its target quickly), or to evade the RES (e.g., to increase the

bioavailability of the minicell). Toxicity is reduced or eliminated because the therapeutic agent is not excreted or processed by the liver and thus does not damage the kidneys or liver, because the minicell-based therapeutic is not activated until entry into the target cell (e.g., in the case of cancer therapeutics or gene therapy). Minicells are of the appropriate size (from about 0.005, 0.1, 0.15 or 0.2 micrometers to about 0.25, 0.3, 0.35, 0.4, 0.45 or 0.5 micrometers) to facilitate deep penetration into the lungs in the cases where administration of the minicell-based therapeutic or diagnostic is via an inhalant (Strong, A. A., et al. 1987. An aerosol generator system for inhalation delivery of pharmacological agents. *Med. Instrum.* 21:189-194). This is due to the fact that minicells can be aerosolized. Without being limited to the following examples, inhalant therapeutic uses of minicells could be applied to the treatment of anaphylactic shock, viral infection, inflammatory reactions, gene therapy for cystic fibrosis, treatment of lung cancers, and fetal distress syndrome.

[0223] Minicells can also display expressed proteins that are enzymes that may have therapeutic and/or diagnostic uses. The enzymes that are displayed may be soluble enzymes that are expressed as fusion proteins with a transmembrane domain of another protein. Display of such enzymes could be used for in vitro assays or for therapeutic benefit.

[0224] Gene therapy applications afforded by minicells generally involve the ability of minicells to deliver DNA to target cells (either for replacement therapy, modification of cell function or to kill cells). Expression plasmids can be delivered to target cells that would encode proteins that could be cytoplasmic or could have intracellular signal sequences that would target the protein to a particular organelle (e.g., mitochondria, nuclei, endoplasmic reticulum, etc.). In the case where minicells are engulfed by the target cell, the minicells themselves could have these intracellular targeting sequences expressed on their surfaces so that the minicells could be 'delivered' to intracellular targets.

[0225] Minicells used for the following therapeutic, discovery, and diagnostic applications can be prepared as described in this application and then stored and/or packaged by a variety of ways, including but not limited to lyophilization, freezing, mixing with preservatives (e.g., antioxidants, glycerol), or otherwise stored and packaged in a fashion similar to methods used for liposome and proteoliposome formulations.

[0226] The small size of minicells (from about 0.005, 0.1, 0.15 or 0.2 micrometers to about 0.25, 0.3, 0.35, 0.4, 0.45 or 0.5 micrometers) makes them suitable for many in vitro diagnostic platforms, including the non-limiting examples of lateral flow, ELISA, HTS, especially those applications requiring microspheres or nanospheres that display many target proteins or other molecules. The use of protoplast or poroplast minicells may be especially useful in this regard. Assay techniques are dependent on cell or particle size, protein (or molecule to be tested) amount displayed on the surface of the cell or particle, and the sensitivity of the assay being measured. In current whole-cell systems, the expression of the protein of interest is limiting, resulting in the higher cell number requirement to satisfy the sensitivity of most assays. However, the relatively large size of cells prevents the incorporation of large numbers of cells in these

assays, e.g. 96, 384, and smaller well formats. In contrast, minicells, protoplasts, and poroplasts are smaller in size and can be manipulated to express high levels of the preselected protein, and can be incorporated into small well assay formats.

#### [0227] I. Types of Minicells

[0228] Minicells are derivatives of cells that lack chromosomal DNA and which are sometimes referred to as anucleate cells. Because eubacterial and archaeobacterial cells, unlike eukaryotic cells, do not have a nucleus (a distinct organelle that contains chromosomes), these non-eukaryotic minicells are more accurately described as being "without chromosomes" or "achromosomal," as opposed to "anucleate." Nonetheless, those skilled in the art often use the term "anucleate" when referring to bacterial minicells in addition to other minicells. Accordingly, in the present disclosure, the term "minicells" encompasses derivatives of eubacterial cells that lack a chromosome; derivatives of archaeobacterial cells that lack their chromosome(s) (Laurence et al., *Nucleoid Structure and Partition in Methanococcus jannaschii: An Archaeon With Multiple Copies of the Chromosome*, *Genetics* 152:1315-1323, 1999); and anucleate derivatives of eukaryotic cells. It is understood, however, that some of the relevant art may use the terms "anucleate minicells" or "anucleate cells" loosely to refer to any of the preceding types of minicells.

#### [0229] I.A. Eubacterial Minicells

[0230] One type of minicell is a eubacterial minicell. For reviews of eubacterial cell cycle and division processes, see Rothfield et al., *Bacterial Cell Division*, *Annu. Rev. Genet.*, 33:423-48, 1999; Jacobs et al., *Bacterial cell division: A moveable feast*, *Proc. Natl. Acad. Sci. USA*, 96:5891-5893, May, 1999; Koch, *The Bacterium's Way for Safe Enlargement and Division*, *Appl. and Envir. Microb.*, Vol. 66, No. 9, pp. 3657-3663; Bouche and Pichoff, *On the birth and fate of bacterial division sites*, *Mol Microbiol.* 29: 19-26; Khachatourians et al., *Cell growth and division in Escherichia coli: a common genetic control involved in cell division and minicell formation*, *J Bacteriol.* 1973. 116: 226-229; Cooper, *The Escherichia coli cell cycle*, *Res Microbiol.* 1990. 141: 17-29; and Danachie and Robinson, "Cell Division: Parameter Values and the Process," in: *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*, Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1987, Volume 2, pages 1578-1592, and references cited therein; and Lutkenhaus et al., "Cell Division," Chapter 101 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2<sup>nd</sup> Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1615-1626, and references cited therein. When DNA replication and/or chromosomal partitioning is altered, membrane-bounded vesicles "pinch off" from parent cells before transfer of chromosomal DNA is completed. As a result of this type of dysfunctional division, minicells are produced which contain an intact outer membrane, inner membrane, cell wall, and all of the cytoplasm components but do not contain chromosomal DNA. See Table 2.

#### [0231] I.B. Eukaryotic Minicells

[0232] The term "eukaryote" is defined as is used in the art, and includes any organism classified as Eucarya that are

usually classified into four kingdoms: plants, animals, fungi and protists. The first three of these correspond to phylogenetically coherent groups. However, the eucaryotic protists do not form a group, but rather are comprised of many phylogenetically disparate groups (including slime molds, multiple groups of algae, and many distinct groups of protozoa). See, e.g., Olsen, G., <http://www.bact.wisc.edu/microtextbook/>. A type of animal of particular interest is a mammal, including, by way of non-limiting example a rat, a mouse, a rabbit, a dog, a cat, a horse, a cow, a pig, a simian and a human.

[0233] Chromosomeless eukaryotic minicells (i.e., anucleate cells) are within the scope of the invention. Platelets are a non-limiting example of eukaryotic minicells. Platelets are anucleate cells with little or no capacity for de novo protein synthesis. The tight regulation of protein synthesis in platelets (Smith et al., Platelets and stroke, *Vasc Med* 4:165-72, 1999) may allow for the over-production of exogenous proteins and, at the same time, under-production of endogenous proteins. Thrombin-activated expression elements such as those that are associated with Bcl-3 (Weyrich et al., Signal-dependent translation of a regulatory protein, Bcl-3, in activated human platelets, *Cell Biology* 95:5556-5561, 1998) may be used to modulate the expression of exogenous genes in platelets.

[0234] As another non-limiting example, eukaryotic minicells are generated from tumor cell lines (Gyongyossy-Issa and Khachatourians, Tumour minicells: single, large vesicles released from cultured mastocytoma cells (1985) *Tissue Cell* 17:801-809; Melton, Cell fusion-induced mouse neuroblastomas HPRT revertants with variant enzyme and elevated HPRT protein levels (1981) *Somatic Cell Genet* 7: 331-344).

[0235] Yeast cells are used to generate fungal minicells. See, e.g., Lee et al., Ibd1p, a possible spindle pole body associated protein, regulates nuclear division and bud separation in *Saccharomyces cerevisiae*, *Biochim Biophys Acta* 3:239-253, 1999; Kopecka et al., A method of isolating anucleated yeast protoplasts unable to synthesize the glucan fibrillar component of the wall *J Gen Microbiol* 81:111-120, 1974; and Yoo et al., Fission yeast Hrp1, a chromodomain ATPase, is required for proper chromosome segregation and its overexpression interferes with chromatin condensation, *Nucl Acids Res* 28:2004-2011, 2000. Cell division in yeast is reviewed by Gould and Simanis, The control of septum formation in fission yeast, *Genes & Dev* 11:2939-51, 1997).

#### [0236] I.C. Archeobacterial Minicells

[0237] The term "archeobacterium" is defined as is used in the art and includes extreme thermophiles and other Archaea. Woese, C. R., L. Magrum. G. Fox. 1978. Archeobacteria. *Journal of Molecular Evolution*. 11:245-252. Three types of Archeobacteria are halophiles, thermophiles and methanogens. By physiological definition, the Archaea (informally, archaees) are single-cell extreme thermophiles (including thermoacidophiles), sulfate reducers, methanogens, and extreme halophiles. The thermophilic members of the Archaea include the most thermophilic organisms cultivated in the laboratory. The aerobic thermophiles are also acidophilic; they oxidize sulfur in their environment to sulfuric acid. The extreme halophiles are aerobic or microaerophilic and include the most salt tolerant organisms known. The sulfate-reducing Archaea reduce sulfate to sul-

fide in extreme environment. Methanogens are strict anaerobes, yet they gave rise to at least two separate aerobic groups: the halophiles and a thermoacidophilic lineage (Olsen, G., <http://www.bact.wisc.edu/microtextbook/>). Non-limiting examples of halophiles include *Halobacterium cutirubrum* and *Halogerax mediterranei*. Non-limiting examples of methanogens include *Methanococcus voltae*; *Methanococcus vanniela*; *Methanobacterium thermoautotrophicum*; *Methanococcus voltae*; *Methanothermus fervidus*; and *Methanosarcina barkeri*. Non-limiting examples of thermophiles include *Azotobacter vinelandii*; *Thermoplasma acidophilum*; *Pyrococcus horikoshii*; *Pyrococcus furiosus*; and Crenarchaeota (extremely thermophilic archaeobacteria) species such as *Sulfolobus solfataricus* and *Sulfolobus acidocaldarius*.

[0238] Archeobacterial minicells are within the scope of the invention. Archeobacteria have homologs of eubacterial minicell genes and proteins, such as the MinD polypeptide from *Pyrococcus furiosus* (Hayashi et al., *EMBO J.* 2001 20:1819-28, Structural and functional studies of MinD ATPase: implications for the molecular recognition of the bacterial cell division apparatus). It is thus possible to create Archeobacterial minicells by methods such as, by way of non-limiting example, overexpressing the product of a min gene isolated from a prokaryote or an archeobacterium; or by disrupting expression of a min gene in an archeobacterium of interest by, e.g., the introduction of mutations thereof or antisense molecules thereto. See, e.g., Laurence et al., Nucleoid Structure and Partition in *Methanococcus jannaschii*: An Archaeon With Multiple Copies of the Chromosome, *Genetics* 152:1315-1323, 1999.

[0239] In one aspect, the invention is drawn to archael minicells. By physiological definition, the Archaea (informally, archaees) are single-cell extreme thermophiles (including thermoacidophiles), sulfate reducers, methanogens, and extreme halophiles. The thermophilic members of the Archaea include the most thermophilic organisms cultivated in the laboratory. The aerobic thermophiles are also acidophilic; they oxidize sulfur in their environment to sulfuric acid. The extreme halophiles are aerobic or microaerophilic and include the most salt tolerant organisms known. The sulfate-reducing Archaea reduce sulfate to sulfide in extreme environment. Methanogens are strict anaerobes, yet they gave rise to at least two separate aerobic groups: the halophiles and a thermoacidophilic lineage (Olsen, G., <http://www.bact.wisc.edu/microtextbook/>).

#### [0240] I.D. Minicells Produced from Diverse Organisms

[0241] There are genes that can be disrupted to cause minicell production that are conserved among the three Kingdoms. For example, SMC (structural maintenance of chromosomes) proteins are conserved among prokaryotes, archeobacteria and eukaryotes (Hirano, SMC-mediated chromosome and mechanics: a conserved scheme from bacteria to vertebrates?, *Genes and Dev.* 13:11-19, 1999; Holmes et al., Closing the ring: Links between SMC proteins and chromosome partitioning, condensation, and supercoiling, *PNAS* 97:1322-1324, 2000; Michiko and Hiranol, *EMBO J.* 17:7139-7148, 1998, ATP-dependent aggregation of single-stranded DNA by a bacterial SMC homodimer, 1998). Mutations in *B. subtilis* smc genes result in the production of minicells (Britton et al., Characterization of a eubacterial smc protein involved in chromosome

partitioning, *Genes and Dev.* 12:1254-1259, 1998; Moriya et al., *A Bacillus subtilis* gene-encoding protein homologous to eukaryotic SMC motor protein is necessary for chromosome partition *Mol Microbiol* 29:179-87, 1998). Disruption of *smc* genes in various cells is predicted to result in minicell production therefrom.

[0242] As another example, mutations in the yeast genes encoding TRF topoisomerases result in the production of minicells, and a human homolog of yeast TRF genes has been stated to exist (Castano et al., A novel family of TRF (DNA topoisomerase I-related function) genes required for proper nuclear segregation, *Nucleic Acids Res* 24:2404-10, 1996). Mutations in a yeast chromodomain ATPase, Hrp1, result in abnormal chromosomal segregation; (Yoo et al., "Fission yeast Hrp1, a chromogomain ATPase, is required for proper chromosome segregation and its overexpression interferes with chromatin condensation," *Nuc. Acids Res.* 28:2004-2001). Disruption of TRF and/or Hrp1 function is predicted to cause minicell production in various cells. Genes involved in septum formation in fission yeast (see, e.g., Gould et al., "The control of septum formation in fission yeast," *Genes and Dev.* 11:2939-2951, 1997) can be used in like fashion.

[0243] As another example, mutations in the *divIVA* gene of *Bacillus subtilis* results in minicell production (Table 2). When expressed in *E. coli* or the yeast *Schizosaccharomyces pombe*, a *B. subtilis* DivIVA-GFP protein is targeted to cell division sites therein, even though clear homologs of DivIVA do not seem to exist in *E. coli* or *S. pombe* (David et al., Promiscuous targeting of *Bacillus subtilis* cell division protein DivIVA to division sites in *Escherichia coli* and fission yeast, *EMBO J.* 19:2719-2727, 2000.) Over- or under-expression of *B. subtilis* DivIVA or a homolog thereof may be used to reduce minicell production in a variety of cells.

#### [0244] II. Production of Minicells

[0245] Eubacterial minicells are produced by parent cells having a mutation in, and/or overexpressing, or under expressing a gene involved in cell division and/or chromosomal partitioning, or from parent cells that have been exposed to certain conditions, that result in aberrant fission of bacterial cells and/or partitioning in abnormal chromosomal segregation during cellular fission (division). The term "parent cells" or "parental cells" refers to the cells from which minicells are produced. Minicells, most of which lack chromosomal DNA (Mulder et al., The *Escherichia coli* *minB* mutation resembles *gyrB* in Defective nucleoid segregation and decreased negative supercoiling of plasmids. *Mol Gen Genet*, 1990, 221: 87-93), are generally, but need not be, smaller than their parent cells. Typically, minicells produced from *E. coli* cells are generally spherical in shape and are about 0.1 to about 0.3  $\mu\text{m}$  in diameter, whereas whole *E. coli* cells are about from about 1 to about 3  $\mu\text{m}$  in diameter and from about about 2 to about 10  $\mu\text{m}$  in length. Micrographs of *E. coli* cells and minicells that have been stained with DAPI (4:6-diamidino-z-phenylindole), a compound that binds to DNA, show that the minicells do not stain while the parent *E. coli* are brightly stained. Such micrographs demonstrate the lack of chromosomal DNA in minicells. (Mulder et al., *Mol. Gen. Genet.* 221:87-93, 1990).

[0246] As shown in Table 2, minicells are produced by several different mechanisms such as, by way of non-

limiting example, the over expression of genes involved in chromosomal replication and partitioning, mutations in such genes, and exposure to various environmental conditions. "Overexpression" refers to the expression of a polypeptide or protein encoded by a DNA introduced into a host cell, wherein the polypeptide or protein is either not normally present in the host cell, or wherein the polypeptide or protein is present in the host cell at a higher level than that normally expressed from the endogenous gene encoding the polypeptide or protein. For example, in *E. coli* cells that overexpress the gene product FtsZ (The FtsZ gene encodes a protein that is involved in regulation of divisions; see Cook and Rothfield, Early stages in development of the *Escherichia coli* cell-division site. *Mol Microbiol*, 1994. 14: p. 485-495; and Lutkenhaus, Regulation of cell division in *E. coli*. *Trends Genet*, 1990. 6: p. 22-25), there is an increase in the formation of minicells (Begg et al., Roles of FtsA and FtsZ in the activation of division sites. *J. Bacteriology*, 1997. 180: 881-884). Minicells are also produced by *E. coli* cells having a mutation in one or more genes of the *min* locus, which is a group of genes that encode proteins that are involved in cell division (de Boer et al., Central role for the *Escherichia coli* *minC* gene product in two different cell division-inhibition systems. *Proc. Natl. Acad. Sci. USA*, 1990. 87: 1129-33; Akerlund et al., Cell division in *Escherichia coli* *minB* mutants. *Mol Microbiol*, 1992. 6: 2073-2083).

[0247] Prokaryotes that have been shown to produce minicells include species of *Escherichia*, *Shigella*, *Bacillus*, *Lactobacillus*, and *Campylobacter*. Bacterial minicell-producing species of particular interest are *E. coli* and *Bacillus subtilis*. *E. coli* is amenable to manipulation by a variety of molecular genetic methods, with a variety of well-characterized expression systems, including many episomal expression systems, factors and elements useful in the present invention. *B. subtilis*, also amenable to genetic manipulation using episomal expression elements, is an important industrial organism involved in the production of many of the world's industrial enzymes (proteases, amylases, etc.), which it efficiently produces and secretes.

[0248] In the case of other eubacterial species, homologs of *E. coli* or *B. subtilis* genes that cause minicell production therein are known or can be identified and characterized as is known in the art. For example, the *min* regions of the chromosome of *Streptococcus pneumoniae* and *Neisseria gonorrhoeae* have been characterized (Massidda et al., Unconventional organization of the division and cell wall gene cluster of *Streptococcus pneumoniae*, *Microbiology* 144:3069-78, 1998; and Ramirez-Arcos et al., *Microbiology* 147:225-237, 2001 and Szeto et al., *Journal of Bacteria* 183(21):6253, 2001, respectively). Those skilled in the art are able to isolate minicell producing (*min*) mutants, or prepare compounds inhibitory to genes that induce a minicell production (e.g., antisense to *min* transcripts).

TABLE 2

Eubacterial Strains, Mutations and Conditions that Promote Minicell Formation			
Species	Strain	Notes	References
<i>Campylobacter jejuni</i>		may occur naturally late in growth cycle	Brock et al., 1987
<i>Bacillus</i>		Mutations in <i>divIVB</i>	Barak et al., 1999

TABLE 2-continued

Eubacterial Strains, Mutations and Conditions that Promote Minicell Formation			
Species	Strain	Notes	References
<i>subtilis</i>		locus (inc. minC, minD ripX mutations)	Sciochetti et al., 1999; Lemon et al., 2001
		smc mutations	Moriya et al., 1998; Britton et al., 1998
		oriC deletions	Moriya et al., 1997; Hassan et al., 1997
		prfA mutations	Pederson and Setlow, 2001
		Mutations in divIVA locus	Cha et al., 1997
	B.s. 168	ts initiation mutation TsB143	Sargent, 1975
<i>Bacillus cereus</i>	WSBC 10030	Induced by exposure to long-chain polyphosphate	Maier et al., 1999
<i>Shigella flexneri</i> (2a)	MC-1		Gemski et al., 1980
<i>S. dysenteriae</i> (1)	MC-V		Gemski et al., 1980
<i>Lactobacillus spp.</i>		Variant minicell-producing strains isolated from grains	Pidoux et al., 1990
<i>Neisseria gonorrhoeae</i>		deletion or overexpression of min homologues	Ramirez-Arcos et al., 2001; Szeto et al., 2001
<i>Escherichia coli</i>		MinA mutations	Frazer et al., 1975; Cohen et al. 1976
		MinB mutations and deletions	Adler et al., 1967; Davie et al., 1984; Schaumberg et al., 1983; Jaffe et al., 1988; Akerlund et al., 1992
	CA8000	cya, crp mutations MukA1 mutation MukE, mukF mutations hns mutation	Kumar et al.; 1979 Hiraaga et al., 1996 Yamanaka et al., 1996
	DS410		Kaidow et al., 1995 Heighway et al., 1989 Curtiss, 1980
	P678-54	$\chi_{1972}$ , $\chi_{1776}$ and $\chi_{2076}$ Temperature-sensitive cell division mutations Induced by over-expression of minB protein	Adler et al. 1967; Allen et al., 1972; Hollenberg et al., 1976 De Boer et al., 1988
		Induced by over-expression of minE protein or derivatives	Pichoff et al., 1995
		Induced by over-production of ftsZ gene	Ward et al., 1985
		Induced by over-expression of sdiA gene	Wang et al., 1991
		Induced by over-expression of min genes from <i>Neisseria gonorrhoeae</i>	Ramirez-Arcos et al., 2001; Szeto et al., 2001
		Induced by exposure to EGTA	Wachi et al., 1999
<i>Legionella pneumophila</i>		Induced by exposure to ampicillin	Elliot et al., 1985

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#### [0283] II.A. Optimized Minicell Construction

[0284] Minicells are produced by several different eubacterial strains and mechanisms including the overexpression of endogenous or exogenous genes involved in cell division, chromosomal replication and partitioning, mutations in such genes, and exposure to various chemical and/or physical conditions. For example, in *E. coli* cells that overexpress the gene product FtsZ (the *ftsZ* gene encodes a protein that is involved in regulation of cell division; see Cook and Rothfield, Early stages in development of the *Escherichia coli* cell-division site. *Mol Microbiol*, 1994. 14: p. 485-495; and Lutkenhaus, Regulation of cell division in *E. coli*. *Trends Genet*, 1990. 6: p. 22-25), there is an increase in the formation of minicells (Begg et al., Roles of FtsA and FtsZ in the activation of division sites. *J. Bacteriology*, 1997. 180: 881-884). Minicells are also produced by *E. coli* cells having a mutation in one or more genes of the *min* locus, which is a group of genes that encode proteins that are involved in cell division (de Boer et al., Central role for the *Escherichia coli* *minC* gene product in two different cell division-inhibition systems. *Proc. Natl. Acad. Sci. USA*, 1990. 87: 1129-33; Akerlund et al., Cell division in *Escherichia coli* *minB* mutants. *Mol Microbiol*, 1992. 6: 2073-2083).

[0285] Eubacterial cells that have been shown to produce minicells include, but are not limited to species of *Escherichia*, *Shigella*, *Bacillus*, *Lactobacillus*, *Legionella* and *Campylobacter*. Bacterial minicell-producing species of particular interest are *E. coli* and *Bacillus subtilis*. These organisms are amenable to manipulation by a variety of molecular and genetic methods, with a variety of well-characterized expression systems, including many episomal and chromosomal expression systems, as well as other factors and elements useful in the present invention.

[0286] The following sections describe genes that may be manipulated so as to stimulate the production of minicells. The invention may include any of these non-limiting examples for the purpose of preparing minicells. Furthermore, these genes and gene products and conditions, may be used in methodologies to identify other gene(s), gene products, biological events, biochemical events, or physiological events that induce or promote the production of minicells. These methodologies include, but are not limited to genetic selection, protein, nucleic acid, or combinatorial chemical library screen, one- or two-hybrid analysis, display selection technologies, e.g. phage or yeast display, hybridization approaches, e.g. array technology, and other high- or low-throughput approaches.

#### [0287] II.A.1. Homologs

[0288] Homologs of these genes and gene products from other organisms may also be used. As used herein, a "homolog" is defined as a nucleic acid or protein having a nucleotide sequence or amino acid sequence, respectively, that is "identical," "essentially identical," "substantially identical," "homologous" or "similar" (as described below) to a

reference sequence which may, by way of non-limiting example, be the sequence of an isolated nucleic acid or protein, or a consensus sequence derived by comparison of two or more related nucleic acids or proteins, or a group of isoforms of a given nucleic acid or protein. Non-limiting examples of types of isoforms include isoforms of differing molecular weight that result from, e.g., alternate RNA splicing or proteolytic cleavage; and isoforms having different post-translational modifications, such as glycosylation; and the like.

[0289] Two sequences are said to be "identical" if the two sequences, when aligned with each other, are exactly the same with no gaps, substitutions, insertions or deletions.

[0290] Two sequences are said to be "essentially identical" if the following criteria are met. Two amino acid sequences are "essentially identical" if the two sequences, when aligned with each other, are exactly the same with no gaps, insertions or deletions, and the sequences have only conservative amino acid substitutions. Conservative amino acid substitutions are as described in Table 3.

TABLE 3

CONSERVATIVE AMINO ACID SUBSTITUTIONS	
Type of Amino Acid Side Chain	Groups of Amino Acids that Are Conservative Substitutions Relative to Each Other
Short side chain	Glycine, Alanine, Serine, Threonine and Methionine
Hydrophobic	Leucine, Isoleucine and Valine
Polar	Glutamine and Asparagine
Acidic	Glutamic Acid and Aspartic Acid
Basic	Arginine, Lysine and Histidine
Aromatic	Phenylalanine, Tryptophan and Tyrosine

[0291] Two nucleotide sequences are "essentially identical" if they encode the identical or essentially identical amino acid sequence. As is known in the art, due to the nature of the genetic code, some amino acids are encoded by several different three base codons, and these codons may thus be substituted for each other without altering the amino acid at that position in an amino acid sequence. In the genetic code, TTA, TTG, CTT, CTC, CTA and CTG encode Leu; AGA, AGG, CGT, CGC, CGA and CGG encode Arg; GCT, GCC, GCA and GCG encode Ala; GGT, GGC, GGA and GGG encode Gly; ACT, ACC, ACA and ACG encode Thr; GTT, GTC, GTA and GTG encode Val; TCT, TCC, TCA and TCG encode Ser; CCT, CCC, CCA and CCG encode Pro; ATA, ATC and ATA encode Ile; GAA and GAG encode Glu; CAA and CAG encode Gln; GAT and GAC encode Asp; AAT and AAC encode Asn; AGT and AGC encode Ser; TAT and TAC encode Tyr; TGT and TGC encode Cys; AAA and AAG encode Lys; CAT and CAC encode His; TTT and TTC encode Phe, TGG encodes Trp; ATG encodes Met; and TGA, TAA and TAG are translation stop codons.

[0292] Two amino acid sequences are "substantially identical" if, when aligned, the two sequences are, (i) less than 30%, preferably  $\leq 20\%$ , more preferably  $\leq 15\%$ , most preferably  $\leq 10\%$ , of the identities of the amino acid residues vary between the two sequences; (ii) the number of gaps between or insertions in, deletions of and/or substitutions of, is  $\leq 10\%$ , more preferably  $\leq 5\%$ , more preferably  $\leq 3\%$ ,



most preferably  $\leq 1\%$ , of the number of amino acid residues that occur over the length of the shortest of two aligned sequences.

[0293] Two sequences are said to be "homologous" if any of the following criteria are met. The term "homolog" includes without limitation orthologs (homologs having genetic similarity as the result of sharing a common ancestor and encoding proteins that have the same function in different species) and paralog (similar to orthologs, yet gene and protein similarity is the result of a gene duplication).

[0294] One indication that nucleotide sequences are homologous is if two nucleic acid molecules hybridize to each other under stringent conditions. Stringent conditions are sequence dependent and will be different in different circumstances. Generally, stringent conditions are selected to be about 5° C. lower than the thermal melting point (T<sub>m</sub>) for the specific sequence at a defined ionic strength and pH. The T<sub>m</sub> is the temperature (under defined ionic strength and pH) at which 50% of the target sequence hybridizes to a perfectly matched probe. Typically, stringent conditions will be those in which the salt concentration is about 0.02 M at pH 7 and the temperature is at least about 60° C.

[0295] Another way by which it can be determined if two sequences are homologous is by using an appropriate algorithm to determine if the above-described criteria for substantially identical sequences are met. Sequence comparisons between two (or more) polynucleotides or polypeptides are typically performed by algorithms such as, for example, the local homology algorithm of Smith and Waterman (Adv. Appl. Math. 2:482, 1981); by the homology alignment algorithm of Needleman and Wunsch (J. Mol. Biol. 48:443, 1970); by the search for similarity method of Pearson and Lipman (Proc. Natl. Acad. Sci. U.S.A. 85:2444, 1988); and by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, version 10.2 Genetics Computer Group (GCG), 575 Science Dr., Madison, Wis.); BLASTP, BLASTN, and FASTA (Altschul et al., J. Mol. Biol. 215:403-410, 1990); or by visual inspection.

[0296] Optimal alignments are found by inserting gaps to maximize the number of matches using the local homology algorithm of Smith and Waterman (1981) Adv. Appl. Math. 2:482-489. "Gap" uses the algorithm of Needleman and Wunsch (1970 J. Mol. Biol. 48:443-453) to find the alignment of two complete sequences that maximizes the number of matches and minimizes the number of gaps. In such algorithms, a "penalty" of about 3.0 to about 20 for each gap, and no penalty for end gaps, is used.

[0297] Homologous proteins also include members of clusters of orthologous groups of proteins (COGs), which are generated by phylogenetic classification of proteins encoded in complete genomes. To date, COGs have been delineated by comparing protein sequences encoded in 43 complete genomes, representing 30 major phylogenetic lineages. Each COG consists of individual proteins or groups of paralogs from at least 3 lineages and thus corresponds to an ancient conserved domain (see Tatusov et al., A genomic perspective on protein families. Science, 278: 631-637, 1997; Tatusov et al., The COG database: new developments in phylogenetic classification of proteins from complete genomes, Nucleic Acids Res. 29:22-28, 2001; Chervitz et al., Comparison of the Complete Sets of Worm and Yeast:

Orthology and Divergence, Science 282:2022-2028, 1998; and <http://www.ncbi.nlm.nih.gov/COG/>).

[0298] The entirety of two sequences may be identical, essentially identical, substantially identical, or homologous to one another, or portions of such sequences may be identical or substantially identical with sequences of similar length in other sequences. In either case, such sequences are similar to each other. Typically, stretches of identical or essentially within similar sequences have a length of  $\geq 12$ , preferably  $\geq 24$ , more preferably  $\geq 48$ , and most preferably  $\geq 96$  residues.

[0299] II.A.2. *Escherichia coli* Genes

[0300] Exemplary genes and gene products from *E. coli* the expression and/or sequence of which can be manipulated so as to stimulate minicell production in *E. coli* or any other organism, as can homologs thereof from any species, include without limitation, the bolA gene (Aldea, M., et al. 1988. Identification, cloning, and expression of bolA, an *ftsZ*-dependent morphogene of *Escherichia coli*. J. Bacteriol. 170:5196-5176; Aldea, M., et al. 1990. Division genes in *Escherichia coli* are expressed coordinately to cell septum requirements by gearbox promoters. EMBO J. 9:3787-3794); the chpA gene (Masuda, Y., et al. 1993. chpA and chpB, *Escherichia coli* chromosomal homologs of the pem locus responsible for stable maintenance of plasmid R100. J. Bacteriol. 175:6850-6856); the chpB gene (Masuda, Y., et al. 1993. chpA and chpB, *Escherichia coli* chromosomal homologs of the pem locus responsible for stable maintenance of plasmid R100. J. Bacteriol. 175:6850-6856); the chpR (chpAI) gene (Masuda, Y., et al. 1993. chpA and chpB, *Escherichia coli* chromosomal homologs of the pem locus responsible for stable maintenance of plasmid R100. J. Bacteriol. 175:6850-6856); the chpS (chpBI) gene (Masuda, Y., et al. 1993. chpA and chpB, *Escherichia coli* chromosomal homologs of the pem locus responsible for stable maintenance of plasmid R100. J. Bacteriol. 175:6850-6856); the crg gene (Redfield, R. J., and A. M. Campbell. 1987. Structure of cryptic lambda prophages. J. Mol. Biol. 198:393-404); the crp gene (Kumar, S., et al. 1979. Control of minicell producing cell division by cAMP-receptor protein complex in *Escherichia coli*. Mol. Gen. Genet. 176:449-450); the cyo gene (Kumar, S., et al. 1979. Control of minicell producing cell division by cAMP-receptor protein complex in *Escherichia coli*. Mol. Gen. Genet. 176:449-450); the dicA gene (Labie, C., et al. 1989. Isolation and mapping of *Escherichia coli* mutations conferring resistance to division inhibition protein DicB. J. Bacteriol. 171:4315-4319); the dicB gene (Labie, C., et al. 1989. Isolation and mapping of *Escherichia coli* mutations conferring resistance to division inhibition protein DicB. J. Bacteriol. 171:4315-4319; Labie, C., et al. 1990. Minicell-forming mutants of *Escherichia coli*: suppression of both DicB- and MinD-dependent division inhibition by inactivation of the minC gene product. J. Bacteriol. 190. 172:5852-5858); the dicC gene (Bejar, S., et al. 1988. Cell division inhibition gene dicB is regulated by a locus similar to lambdaoid bacteriophage immunity loci. Mol. Gen. Genet. 212:11-19); the dif gene (Tetart, F., and J. P. Bouche. 1992. Regulation of the expression of the cell-cycle gene *ftsZ* by DicF antisense RNA. Division does not require a fixed number of FtsZ molecules. Mol. Microbiol. 6:615-620); the dif gene (Kuempe, P. L., et al. 1991. dif, a recA-independent recombination site in the terminus region of the chromosome of *Escheri-*

*chia coli*. New Biol. 3:799-811); the *dksA* gene (Yamanaka, K., et al. 1994. Cloning, sequencing, and characterization of multicopy suppressors of a *mukB* mutation in *Escherichia coli*. Mol. Microbiol. 13:301-312); the *dnaK* gene (Paek, K. H., and G. C. Walker. 1987. *Escherichia coli* *dnaK* null mutants are inviable at high temperature. J. Bacteriol. 169:283-290); the *dnaJ* gene (Hoffman, H. J., et al. 1992. Activity of the Hsp70 chaperone complex—DnaK, DnaJ, and GrpE—in initiating phage lambda DNA replication by sequestering and releasing lambda P protein. Proc. Natl. Acad. Sci. 89:12108-12111); the *fcsA* gene (Kudo, T., et al. 1977. Characteristics of a cold-sensitive cell division mutant *Escherichia coli* K-12. Agric. Biol. Chem. 41:97-107); the *fic* gene (Utsumi, R., et al. 1982. Involvement of cyclic AMP and its receptor protein in filamentation of an *Escherichia coli* *fic* mutant. J. Bacteriol. 151:807-812; Komano, T., et al. 1991. Functional analysis of the *fic* gene involved in regulation of cell division. Res. Microbiol. 142:269-277); the *fis* gene (Spaeny-Dekking, L. et al. 1995. Effects of N-terminal deletions of the *Escherichia coli* protein Fis on the growth rate, TRNA (2Ser) expression and cell morphology. Mol. Gen. Genet. 246:259-265); the *ftsA* gene (Bi, E., and J. Lutkenhaus. 1990. Analysis of *ftsZ* mutations that confer resistance to the cell division inhibitor SulA (SfiA). J. Bacteriol. 172:5602-5609; Dai, K., and J. Lutkenhaus. 1992. The proper ratio of FtsZ to FtsA is required for cell division to occur in *Escherichia coli*. J. Bacteriol. 174:6145-6151); the *ftsE* gene (Taschner, P. E. et al. 1988. Division behavior and shape changes in isogenic *ftsZ*, *ftsQ*, *ftsA*, *pbpB*, and *ftsE* cell division mutants of *Escherichia coli* during temperature shift experiments. J. Bacteriol. 170:1533-1540); the *ftsH* gene (Ogura, T. et al. 1991. Structure and function of the *ftsH* gene in *Escherichia coli*. Res. Microbiol. 142:279-282); the *ftsI* gene (Begg, K. J., and W. D. Donachie. 1985. Cell shape and division in *Escherichia coli*: experiments with shape and division mutants. J. Bacteriol. 163:615-622); the *ftsJ* gene (Ogura, T. et al. 1991. Structure and function of the *ftsH* gene in *Escherichia coli*. Res. Microbiol. 142:279-282); the *ftsL* gene (Guzman, et al. 1992. FtsL, an essential cytoplasmic membrane protein involved in cell division in *Escherichia coli*. J. Bacteriol. 174:7716-7728); the *ftsN* gene (Dai, K. et al. 1993. Cloning and characterization of *ftsN*, an essential cell division gene in *Escherichia coli* isolated as a multicopy suppressor of *ftsA12*(Ts). J. Bacteriol. 175:3790-3797); the *ftsQ* gene (Wang, X. D. et al. 1991. A factor that positively regulates cell division by activating transcription of the major cluster of essential cell division genes of *Escherichia coli*. EMBO J. 10:3362-3372); the *ftsW* gene (Khattar, M. M. et al. 1994. Identification of FtsW and characterization of a new *ftsW* division mutant of *Escherichia coli*. J. Bacteriol. 176:7140-7147); the *ftsX* (*ftsS*) gene (Salmond, G. P. and S. Plakidou. 1984. Genetic analysis of essential genes in the *ftsE* region of the *Escherichia coli* genetic map and identification of a new cell division gene, *ftsS*. Mol. Gen. Genet. 197:304-308); the *ftsY* gene (Gill, D. R. and G. P. Salmond. 1990. The identification of the *Escherichia coli* *ftsY* gene product: an unusual protein. Mol. Microbiol. 4:575-583); the *ftsZ* gene (Ward, J. E., and J. Lutkenhaus. 1985. Overproduction of FtsZ induces minicell formation. Cell. 42:941-949; Bi, E., and J. Lutkenhaus. 1993. Cell division inhibitors SulA and MinCD prevent formation of the FtsZ ring. J. Bacteriol. 175:1118-1125); the *gyrB* gene (Mulder, E., et al. 1990. The *Escherichia coli* *minB* mutation resembles *gyrB* in defective nucleoid segregation and decreased negative supercoiling of plasmids. Mol. Gen. Genet. 221:87-93); the *hlfB* (*ftsH*) gene (Herman, C., et al. 1993. Cell growth and lambda phage development controlled by the same essential *Escherichia coli* gene, *ftsH/hlfB*. Proc. Natl. Acad. Sci. 90:10861-10865); the *hfq* gene (Takada, A., et al. 1999. Negative regulatory role of the *Escherichia coli* *hfq* gene in cell division. Biochem. Biophys. Res. Commun. 266:579-583); the *hipA* gene (Scherrer, R., and H. S. Moyed. 1988. Conditional impairment of cell division and altered lethality in *hipA* mutants of *Escherichia coli* K-12. J. Bacteriol. 170:3321-3326); the *hipB* gene (Hendricks, E. C., et al. 2000. Cell division, guillotining of dimer chromosomes and SOS induction in resolution mutants (*dif*, *xerC* and *xerD*) of *Escherichia coli*. Mol. Microbiol. 36:973-981); the *hns* gene (Kaidow, A., et al. 1995. Anucleate cell production by *Escherichia coli* delta *hns* mutant lacking a histone-like protein, H-NS. J. Bacteriol. 177:3589-3592); the *htrB* gene (Karow, M., et al. 1991. Complex phenotypes of null mutations in the *htr* genes, whole products are essential for *Escherichia coli* growth at elevated temperatures. Res. Microbiol. 142:289-294); the *lpxC* (*envA*) gene (Beall, B., and J. Lutkenhaus. 1987. Sequence analysis, transcriptional organization, and insertional mutagenesis of the *envA* gene of *Escherichia coli*. J. Bacteriol. 169:5408-5415; Young, K., et al. 1995. The *envA* permeability/cell division gene of *Escherichia coli* encodes the second enzyme of lipid A biosynthesis. UDP-3-O-(R-3-hydroxymyristoyl)-N-acetylglucosamine deacetylase. J. Biol. Chem. 270:30384-30391); the *malE* gene (Pichoff, S., et al. 1997. MinCD-independent inhibition of cell division by a protein that fuses MalE to the topological specificity factor MinE. J. Bacteriol. 179:4616-4619); the *minA* gene (Davie, E., et al. 1984. Genetic basis of minicell formation in *Escherichia coli* K-12. J. Bacteriol. 158:1202-1203); the *minB* gene (Davie, E., et al. 1984. Genetic basis of minicell formation in *Escherichia coli* K-12. J. Bacteriol. 158:1202-1203); the *minC* gene (de Boer, P. A., et al. 1990. Central role for the *Escherichia coli* *minC* gene product in two different cell division-inhibition systems. Proc. Natl. Acad. Sci. 87:1129-1133); the *minD* gene (Lapie, C., et al. 1990. Minicell-forming mutants of *Escherichia coli*: suppression of both *DicB*- and *MinD*-dependent division inhibition by inactivation of the *minC* gene product. J. Bacteriol. 172:5852-5855; Hayashi, I., et al. 2001. Structural and functional studies of MinD ATPase: implications for the molecular recognition of the bacterial cell division apparatus. EMBO J. 20:1819-1828); the *minE* gene (de Boer, P. A., et al. 1989. A division inhibitor and a topological specificity factor coded for by the minicell locus determine proper placement of the division septum in *E. coli*. Cell. 56:641-649); the *mreB* gene (Doi, M., et al. 1988. Determinations of the DNA sequence of the *mreB* gene and of the gene products of the *mre* region that function in formation of the rod shape of *Escherichia coli* cells. J. Bacteriol. 170:4619-4624); the *mreC* gene (Wachi, M., et al. 1989. New *mre* genes *mreC* and *mreD*, responsible for formation of the rod shape of *Escherichia coli* cells. J. Bacteriol. 171:6511-6516); the *mreD* gene (Wachi, M., et al. 1989. New *mre* genes *mreC* and *mreD*, responsible for formation of the rod shape of *Escherichia coli* cells. J. Bacteriol. 171:6511-6516); the *mukA* gene (Hiraga, S., et al. 1989. Chromosome partitioning in *Escherichia coli*: novel mutants producing anucleate cells. J. Bacteriol. 171:1496-1505); the *mukB* gene (Hiraga, S., et al. 1991. Mutants defective in

- chromosome partitioning in *E. coli*. Res. Microbiol. 142:189-194); the mukE gene (Yamanaka, K., et al. 1996. Identification of two new genes, mukE and mukF, involved in chromosome partitioning in *Escherichia coli*. Mol. Gen. Genet. 250:241-251; Yamazoe, M., et al. 1999. Complex formation of MukB, MukE and MukF proteins involved in chromosome partitioning in *Escherichia coli*. EMBO J. 18:5873-5884); the mukF gene (Yamanaka, K., et al. 1996. Identification of two new genes, mukE and mukF, involved in chromosome partitioning in *Escherichia coli*. Mol. Gen. Genet. 250:241-251; Yamazoe, M., et al. 1999. Complex formation of MukB, MukE and MukF proteins involved in chromosome partitioning in *Escherichia coli*. EMBO J. 18:5873-5884); the parC gene (Kato, J., et al. 1988. Gene organization in the region containing a new gene involved in chromosome partition in *Escherichia coli*. J. Bacteriol. 170:3967-3977); the parE gene (Roberts, R. C., et al. 1994. The parDE operon of the broad-host-range plasmid RK2 specifies growth inhibition associated with plasmid loss. J. Mol. Biol. 237:35-51); the pbpA gene (Rodriguez, M. C., and M. A. de Pedro. 1990. Initiation of growth in pbpA<sup>ts</sup> and rodA<sup>ts</sup> mutants of *Escherichia coli*. FEMS Microbiol. Lett. 60:235-239); the pcnB gene (Makise, M., et al. 1999. Identification of a high-copy-number plasmid suppressor of a lethal phenotype caused by mutant DnaA protein which has decreased intrinsic ATPase activity. Biol. Pharm. Bull. 22:904-909); the parF (plsC in *E. coli*) gene product from *Salmonella* (Luttinger, A. L., et al. 1991. A cluster of genes that affects nucleoid segregation in *Salmonella typhimurium*. New Biol. 3:687-697); the rpoS gene (Cam, K., et al. 1995. Sigma S-dependent overexpression of ftsZ in an *Escherichia coli* K-12 rpoB mutant that is resistant to the division inhibitors DicB and DicF RNA. Mol. Gen. Genet. 248:190-194); the rcsB gene (Gervais, F. G., et al. 1992. The rcsB gene, a positive regulator of colanic acid biosynthesis in *Escherichia coli*, is also an activator of ftsZ expression. J. Bacteriol. 174:3964-3971); the rcsF gene (Gervais, F. G., and G. R. Drapeau. 1992. Identification, cloning, and characterization of rcsF, a new regulator gene for exopolysaccharide synthesis that suppresses the division mutation ftsZ84 in *Escherichia coli* K-12. J. Bacteriol. 174:8016-8022); the rodA gene (Rodriguez, M. C., and M. A. de Pedro. 1990. Initiation of growth in pbpA<sup>ts</sup> and rodA<sup>ts</sup> mutants of *Escherichia coli*. FEMS Microbiol. Lett. 60:235-239); the sdiA (sulB, sfiB) gene (Wang, X. D., et al. 1991. A factor that positively regulates cell division by activating transcription of the major cluster of essential cell division genes of *Escherichia coli*. EMBO J. 10:3363-3372); the sefA (fabZ) gene (Mohan, S., et al. 1994. An *Escherichia coli* gene (FabZ) encoding (3R)-hydroxymyristoyl acyl carrier protein dehydrase. Relation to fabA and suppression of mutations in lipid A biosynthesis. J. Biol. Chem. 269:32896-32903); the sfiC gene (D'Ari, R., and O. Huisman. 1983. Novel mechanism of cell division inhibition associated with the SOS response in *Escherichia coli*. J. Bacteriol. 156:243-250); the sulA gene (Bi, E., and J. Lutkenhaus. 1990. Interaction between the min locus and ftsZ. J. Bacteriol. 172:5610-5616; Bi, E., and J. Lutkenhaus. 1993. Cell division inhibitors SulA and MinCD prevent formation of the FtsZ ring. J. Bacteriol. 175:1118-1125); the stfZ gene (Dewar, S. J., and W. D. Donachie. 1993. Antisense transcription of the ftsZ-ftsA gene junction inhibits cell division in *Escherichia coli*. J. Bacteriol. 175:7097-7101); the tolC gene (Hiraga, S., et al. 1989. Chromosome partitioning in *Escherichia coli*: novel mutants producing anucleate cells. J. Bacteriol. 171:1496-1505; Hiraga, S., et al. 1991. Mutants defective in chromosome partitioning in *E. coli*. Res. Microbiol. 142:189-194); and the zipA gene (Hale, C. A., and P. A. de Boer. 1997. Direct binding of FtsZ to ZipA, an essential component of the septal ring structure that mediates cell division in *E. coli*. Cell. 88:175-185).
- [0301] The guanosine 5'-diphosphate 3' diphosphate (ppGpp) or guanosine 5'-triphosphate 3' diphosphate (pppGpp) nucleotides, collectively (p)ppGpp, found in *E. coli* or in other members of the Eubacteria, Eucarya or Archaea may be employed to produce minicells (Vinella, D., et al. 1993. Penicillin-binding protein 2 inactivation in *Escherichia coli* results in cell division inhibition, which is relieved by FtsZ overexpression. J. Bacteriol. 175:6704-6710; Navarro, F., et al. Analysis of the effect of ppGpp on the ftsQAZ operon in *Escherichia coli*. Mol. Microbiol. 29:815-823). The levels, or rate of production of (p)ppGpp may be increased or decreased. By way of non-limiting example, increased (p)ppGpp production results from induction of the stringent response. The stringent response in *E. coli* is a physiological response elicited by a failure of the capacity for tRNA aminoacylation to keep up with the demands of protein synthesis. This response can be provoked either by limiting the availability of amino acids or by limiting the ability to aminoacylate tRNA even in the presence of abundant cognate amino acids. Many features of the stringent response behave as if they are mediated by accumulation of (p)ppGpp. The accumulation of (p)ppGpp can also be provoked by nutritional or other stress conditions in addition to a deficiency of aminoacyl-tRNA. See Cashel et al., "The Stringent Response," Chapter 92 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 1, pages 1458-1496, and references cited therein.
- [0302] By way of non-limiting example, factors that may provoke the stringent response include the *lyt* gene or gene product (Harkness, R. E., et al. 1992. Genetic mapping of the *lytA* and *lytB* loci of *Escherichia coli*, which are involved in penicillin tolerance and control of the stringent response. Can J. Microbiol. 38:975-978), the *relA* gene or gene product (Vinella, D., and R. D'Ari. 1994. Thermoinducible filamentation in *Escherichia coli* due to an altered RNA polymerase beta subunit is suppressed by high levels of ppGpp. J. Bacteriol. 176:96-972), the *relB* gene or gene product (Christensen, S. K., et al. 2001. RelE, a global inhibitor of translation, is activated during nutritional stress. Proc. Natl. Acad. Sci. 98:14328-14333), the *relC* (*rplK*) gene or gene product (Yang, X., and E. E. Ishiguro. 2001. Involvement of the N Terminus of Ribosomal Protein L11 in Regulation of the RelA Protein of *Escherichia coli*. J. Bacteriol. 183:6532-6537), the *relX* gene or gene product (St. John, A. C., and A. L. Goldberg. 1980. Effects of starvation for potassium and other inorganic ions on protein degradation and ribonucleic acid synthesis in *Escherichia coli*. J. Bacteriol. 143:1223-1233), the *spoT* gene or gene product (Vinella, D., et al. 1996. Mecillinam resistance in *Escherichia coli* is conferred by loss of a second activity of the AroK protein. J. Bacteriol. 178:3818-3828), the *gpp* gene or gene product (Keasling, J. D., et al. 1993. Guanosine pentaphosphate phosphohydrolase of *Escherichia coli* is a long-chain exopolyphosphatase. Proc. Natl. Acad. Sci.

90:7029-7033), the *ndk* gene or gene product (Kim, H. Y., et al. 1998. Alginate, inorganic polyphosphate, GTP and ppGpp synthesis co-regulated in *Pseudomonas aeruginosa*: implications for stationary phase survival and synthesis of RNA/DNA precursors. *Mol. Microbiol.* 27:717-725), the *rpoB* gene or gene product (Vinella, D., and R. D'Ari. 1994. Thermoinducible filamentation in *Escherichia coli* due to an altered RNA polymerase beta subunit is suppressed by high levels of ppGpp. *J. Bacteriol.* 176:96-972), the *rpoC* gene or gene product (Bartlett, M. S., et al. 1998. RNA polymerase mutants that destabilize RNA polymerase-promoter complexes alter NTP-sensing by *rrn* P1 promoters. *J. Mol. Biol.* 279:331-345), the *rpoD* gene or gene product (Hernandez, V. J., and M. Cashel. 1995. Changes in conserved region 3 of *Escherichia coli* sigma 70 mediate ppGpp-dependent functions in vivo. 252:536-549), *glnF* gene or gene product (Powell, B. S., and D. L. Court. 1998. Control of *ftsZ* expression, cell division, and glutamine metabolism in Luria-Bertani medium by the alarmone ppGpp in *Escherichia coli*. *J. Bacteriol.* 180:1053-1062), or *glnD* gene or gene product (Powell, B. S., and D. L. Court. 1998. Control of *ftsZ* expression, cell division, and glutamine metabolism in Luria-Bertani medium by the alarmone ppGpp in *Escherichia coli*. *J. Bacteriol.* 180:1053-1062). These genes or gene products, and/or expression thereof, may be manipulated to create minicells.

#### [0303] II.A.3. *Bacillus subtilis* Genes

[0304] Exemplary genes and gene products from *B. subtilis*, the expression and/or sequence of which can be manipulated so as to stimulate minicell production in *B. subtilis* or any other organism, as can homologs thereof from any species, include without limitation, the *divI* (*divD*) gene (Van Alstyne, D., and M. I. Simon. 1971. Division mutants of *Bacillus subtilis*: isolation of PBS1 transduction of division-specific markers. *J. Bacteriol.* 108:1366-1379); the *divIB* (*dds*, *ftsQ*) gene (Harry, E. J., et al. 1993. Characterization of mutations in *divIB* of *Bacillus subtilis* and cellular localization of the DivIB protein. *Mol. Microbiol.* 7:611-621; Harry E. J., et al. 1994. Expression of *divIB* of *Bacillus subtilis* during vegetative growth. *J. Bacteriol.* 176:1172-1179); the *divIC* gene product from *B. subtilis* or homologs of this gene or gene product found in other members of the Eubacteria, Eucarya or Archaea may be employed to produce minicells (Levin, P. A., and R. Losick. 1994. Characterization of a cell division gene from *Bacillus subtilis* that is required for vegetative and sporulation septum formation. *J. Bacteriol.* 176:1451-1459; Katis, V. L., et al. 1997. The *Bacillus subtilis* division protein DivIC is a highly abundant membrane-bound protein that localizes to the division site; the *divII* (*divC*) gene (Van Alstyne, D., and M. I. Simon. 1971. Division mutations of *Bacillus subtilis*: isolation and PBS1 transduction of division-specific markers. *J. Bacteriol.* 108:1366-1379); the *divIVA* (*divD*) gene (Cha, J. -H., and G. C. Stewart. 1997. The *divIVA* minicell locus of *Bacillus subtilis*. *J. Bacteriol.* 179:1671-1683); the *divIVC* (*divA*) gene (Van Alstyne, D., and M. I. Simon. 1971. Division mutations of *Bacillus subtilis*: isolation and PBS1 transduction of division-specific markers. *J. Bacteriol.* 108:1366-1379); the *divV* (*divB*) gene (Van Alstyne, D., and M. I. Simon. 1971. Division mutations of *Bacillus subtilis*: isolation and PBS1 transduction of division-specific markers. *J. Bacteriol.* 108:1366-1379); the *erzA* (*ytwP*) gene (Levin, P. A., et al. 1999. Identification and regulation of a negative regulator of FtsZ ring formation in *Bacillus subti-*

*lis*. *Proc. Natl. Acad. Sci.* 96:9642-9647); the *ftsA* (*spoIIN*) gene (Feucht, A., et al. 2001. Cytological and biochemical characterization of the FtsA cell division protein of *Bacillus subtilis*. *Mol. Microbiol.* 40:115-125); the *ftsE* gene (Yoshida, K., et al. 1994. Cloning and nucleotide sequencing of a 15 kb region of the *Bacillus subtilis* genome containing the *iol* operon. *Microbiology.* 140:2289-2298); the *ftsH* gene (Deuerling, E., et al. 1995. The *ftsH* gene of *Bacillus subtilis* is transiently induced after osmotic and temperature upshift. *J. Bacteriol.* 177:4105-4112; Wehrl, W., et al. 2000. The FtsH protein accumulates at the septum of *Bacillus subtilis* during cell division and sporulation. *J. Bacteriol.* 182:3870-3873); the *ftsK* gene (Sciochetti, S. A., et al. 2001. Identification and characterization of the *dif* Site from *Bacillus subtilis*. *J. Bacteriol.* 183:1058-1068); the *ftsL* (*yiiD*) gene (Daniel, R. A., et al. 1998. Characterization of the essential cell division gene *ftsL* (*yiiD*) of *Bacillus subtilis* and its role in the assembly of the division apparatus. *Mol. Microbiol.* 29:593-604); the *ftsW* gene (Ikeda, M., et al. 1989. Structural similarity among *Escherichia coli* FtsW and RodA proteins and *Bacillus subtilis* SpoVE protein, which function in cell division, cell elongation, and spore formation, respectively. *J. Bacteriol.* 171:6375-6378); the *ftsX* gene (Reizer, J., et al. 1998. A novel protein kinase that controls carbon catabolite repression in bacteria. *Mol. Microbiol.* 27:1157-1169); the *ftsZ* gene (Beall, B., and J. Lutkenhaus). FtsZ in *Bacillus subtilis* is required for vegetative septation and for asymmetric septation during sporulation. *Genes and Dev.* 5:447-45); the *gcaD* gene (Hove-Jensen, B. 1992. Identification of *tms-26* as an allele of the *gcaD* gene, which encodes N-acetylglucosamine 1-phosphate uridylyltransferase in *Bacillus subtilis*. *J. Bacteriol.* 174:6852-6856); the *gid* (*glyC*) gene (Kunst, F., et al. 1997. The complete genome sequence of the gram-positive bacterium *Bacillus subtilis*. *Nature.* 390:237-238); the *gidA* gene (Ogasawara, N., and H. Yoshikawa. 1992. Genes and their organization in the replication origin region of the bacterial chromosome. *Mol. Microbiol.* 6:629-634; Nakayashiki, T., and H. Inokuchi. 1998. Novel temperature-sensitive mutants of *Escherichia coli* that are unable to grow in the absence of wild-type tRNA<sup>6Leu</sup>. *J. Bacteriol.* 180:2931-2935); the *gidB* gene (Ogasawara, N., and H. Yoshikawa. 1992. Genes and their organization in the replication origin region of the bacterial chromosome. *Mol. Microbiol.* 6:629-634; Nakayashiki, T., and H. Inokuchi. 1998. Novel temperature-sensitive mutants of *Escherichia coli* that are unable to grow in the absence of wild-type tRNA<sup>6Leu</sup>. *J. Bacteriol.* 180:2931-2935); the *lytC* (*cwlB*) gene (Blackman, S. A., et al. 1998. The role of autolysins during vegetative growth of *Bacillus subtilis* 168. *Microbiology.* 144:73-82); the *lytD* (*cwlG*) gene (Blackman, S. A., et al. 1998. The role of autolysins during vegetative growth of *Bacillus subtilis* 168. *Microbiology.* 144:73-82); the *lytE* (*cwlF*) gene (Ishikawa, S., et al. 1998. Regulation of a new cell wall hydrolase gene, *cwlF*, which affects cell separation in *Bacillus subtilis*. *J. Bacteriol.* 180:23549-2555); the *lytF* (*cwlE*, *yhdD*) gene (Ohnishi, R., et al. 1999. Peptidoglycan hydrolase *lytF* plays a role in cell separation with *CwlF* during vegetative growth of *Bacillus subtilis*. *J. Bacteriol.* 181:3178-1384); the *maf* gene (Butler, Y. X., et al. 1993. Amplification of the *Bacillus subtilis* *maf* gene results in arrested septum formation. *J. Bacteriol.* 175:3139-3145); the *minC* gene (Varley, A. W., and G. C. Stewart. 1992. The *divIVB* region of the *Bacillus subtilis* chromosome encodes homologs of *Escherichia coli* septum placement (*minCD*)

- and cell shape (mreBCD) determinants. *J. Bacteriol.* 174:6729-6742; Barak, I., et al. 1998. MinCD proteins control the septation process during sporulation of *Bacillus subtilis*. *J. Bacteriol.* 180:5327-5333; the minD gene (Varley, A. W., and G. C. Stewart. 1992. The divIVB region of the *Bacillus subtilis* chromosome encodes homologs of *Escherichia coli* septum placement (minCD) and cell shape (mreBCD) determinants. *J. Bacteriol.* 174:6729-6742; Barak, I., et al. 1998. MinCD proteins control the septation process during sporulation of *Bacillus subtilis*. *J. Bacteriol.* 180:5327-5333); the pbpB gene (Daniel, R. A., and J. Errington. 2000. Intrinsic instability of the essential cell division protein FtsL of *Bacillus subtilis* and a role for DivIB protein in FtsL turnover. *Mol. Microbiol.* 35:278-289); the ponA gene (Pederson, L. B., et al. Septal localization of penicillin-binding protein 1 in *Bacillus subtilis*. *J. Bacteriol.* 181:3201-3211); the prfA gene (Popham, D. L., and P. Setlow. 1995. Cloning, nucleotide sequence, and mutagenesis of the *Bacillus subtilis* ponA operon, which codes for penicillin-binding protein (PBP) 1 and a PBP-related factor. *J. Bacteriol.* 177:326-335); the rodB gene (Burdett, I. D. 1979. Electron microscope study of the rod-to-coccus shape change in a temperature-sensitive rod-mutant of *Bacillus subtilis*. *J. Bacteriol.* 137:1395-1405; Burdett, I. D. 1980. Quantitative studies of rod-coccus morphogenesis in a temperature-sensitive rod-mutant of *Bacillus subtilis*. *J. Gen. Microbiol.* 121:93-103); the secA gene (Sadaie, Y., et al. 1991. Sequencing reveals similarity of the wild-type div+ gene of *Bacillus subtilis* to the *Escherichia coli* secA gene. *Gene.* 98:101-105); the smc gene (Britton, R. A., et al. 1998. Characterization of a prokaryotic SMC protein involved in chromosome partitioning. *Genes Dev.* 12:1254-1259; Moriya, S., et al. 1998. A *Bacillus subtilis* gene-encoding protein homologous to eukaryotic SMC motor protein is necessary for chromosome partition. *Mol. Microbiol.* 29:179-187; Hirano, M., and T. Hirano. 1998. ATP-dependent aggregation of single-stranded DNA by a bacterial SMC homodimer. *EMBO J.* 17:7139-7148); the spoIIIE gene (Feucht, a., et al. 1996. Bifunctional protein required for asymmetric cell division and cell-specific transcription in *Bacillus subtilis*. *Genes Dev.* 10:794-803; Khvorova, A., et al. 1998. The spoIIIE locus is involved in the Spo0A-dependent switch in the localization of FtsZ rings in *Bacillus subtilis*. *J. Bacteriol.* 180:1256-1260; Lucet, I., et al. 2000. Direct interaction between the cell division protein FtsZ and the cell differentiation protein SpoIIIE. *EMBO J.* 19:1467-1475); the spo0A gene (Ireton, K., et al. 1994. spo0J is required for normal chromosome segregation as well as the initiation of sporulation in *Bacillus subtilis*. *J. Bacteriol.* 176:5320-5329); the spoIVF gene (Lee, S., and C. W. Price. 1993. The minCD locus of *Bacillus subtilis* lacks the minE determinant that provides topological specificity to cell division. *Mol. Microbiol.* 7:601-610); the spo0J gene (Lin, D. C., et al. 1997. Bipolar localization of a chromosome partition protein in *Bacillus subtilis*. *Proc. Natl. Acad. Sci.* 94:4721-4726; Yamaichi, Y., and H. Niki. 2000. Active segregation by the *Bacillus subtilis* partitioning system in *Escherichia coli*. *Proc. Natl. Acad. Sci.* 97:14656-14661); the smc gene (Moriya, S., et al. 1998. A *Bacillus subtilis* gene-encoding protein homologous to eukaryotic SMC motor protein is necessary for chromosome partition. *Mol. Microbiol.* 29:179-187); the ripX gene (ciochetti, S. A. et al. 1999. The ripX locus of *Bacillus subtilis* encodes a site-specific recombinase involved in proper chromosome partitioning. *J. Bacteriol.* 181:6053-6062); and the spoIIIE gene (Wu, L. J., and J. Errington. 1994. *Bacillus subtilis* spoIIIE protein required for DNA segregation during asymmetric cell division. *Science.* 264:572-575); the gene corresponding to the *B. subtilis* mutant allele ts-31 (Errington, J., and A. D. Richard. Cell division during growth and sporulation. In A. L. Sonenshein, J. A. Hoch., and R. Losick (eds.). *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); the gene corresponding to the *B. subtilis* mutant allele ts-526 (Id.); the yacA gene (Kunst, F., et al. 1997. The complete genome sequence of the gram-positive bacterium *Bacillus subtilis*. *Nature.* 390:237-238); the yfhF gene (Kunst, F., et al. 1997. The complete genome sequence of the gram-positive bacterium *Bacillus subtilis*. *Nature.* 390:237-238); the yfhK gene (Kunst, F., et al. 1997. The complete genome sequence of the gram-positive bacterium *Bacillus subtilis*. *Nature.* 390:237-238); the yjoB gene (Kunst, F., et al. 1997. The complete genome sequence of the gram-positive bacterium *Bacillus subtilis*. *Nature.* 390:237-238); and the ywbG gene (Smith, T. J., et al. 2000. Autolysins of *Bacillus subtilis*: multiple enzymes with multiple functions. *Microbiology.* 146:249-262).
- [0305]** II.A.3. *Saccharomyces cerevisiae* Genes
- [0306]** Exemplary genes and gene products from *S. cerevisiae* the expression and/or sequence of which can be manipulated so as to stimulate minicell production in any organism, as can homologs thereof from any species, include without limitation, the trf gene product family (TRF1, TRF2, TRF3, TRF4, and TRF5) from *Saccharomyces cerevisiae* (Sadoff, B. U., et al. 1995. Isolation of mutants of *Saccharomyces cerevisiae* requiring DNA topoisomerase I. *Genetics.* 141:465-479; Castano, I. B., et al. 1996. A novel family of TRF (DNA topoisomerase I-related function) genes required for proper nuclear segregation. *Nucleic Acids Res.* 24:2404-2410); the 1BD1 gene product from *Saccharomyces cerevisiae* (Lee, J., et al. 1999. Ibd1p, a possible spindle pole body associated protein, regulates nuclear division and bud separation in *Saccharomyces cerevisiae*. *Biochim. Biophys. Acta.* 1449:239-253); the plo1 gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics.* 155:1541-1534); the cdc7 locus product(s) from *Saccharomyces cerevisiae* or homologues of this found in other members of the Eubacteria, Eucarya or Archaea may be employed to produce minicells (Biggins, S. et al. 2001. Genes involved in sister chromatid separation and segregation in the budding yeast *Saccharomyces cerevisiae*. *Genetics.* 159:453-470); the cdc15 locus product(s) from *Saccharomyces cerevisiae* or homologues of this found in other members of the Eubacteria, Eucarya or Archaea may be employed to produce minicells (Mah, A. S., et al. 2001. Protein kinase Cdc15 activates the Dbf2-Mob1 kinase complex. *Proc. Natl. Acad. Sci.* 98:7325-7330); the cdc11 locus product(s) from *Saccharomyces cerevisiae* or homologues of this found in other members of the Eubacteria, Eucarya or Archaea may be employed to produce minicells (Fares, H., et al. 1996. Identification of a developmentally regulated septin and involvement of the septins in spore formation in *Saccharomyces cerevisiae*. *J. Cell Biol.* 132:399-411); the spg1 locus product(s) from *Saccharomyces cerevisiae* or homologues of this found in other members of the Eubac-

teria, Eucarya or Archaea may be employed to produce minicells (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *sid2* locus product(s) from *Saccharomyces cerevisiae* or homologues of this found in other members of the Eubacteria, Eucarya or Archaea may be employed to produce minicells (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *cdc8* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *rho1* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *mpd1* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *mpd2* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *smv2* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *cdc16* gene product from *Saccharomyces cerevisiae* (Heichman, K. A., and J. M. Roberts. 1996. The yeast CDC16 and CDC27 genes restrict DNA replication to once per cell cycle. *Cell*. 85:39-48); the *dma1* gene product from *Saccharomyces cerevisiae* (Murone, M., and V. Simanis. 1996. The fission yeast *dma1* gene is a component of the spindle assembly checkpoint, required to prevent septum formation and premature exit from mitosis if spindle function is compromised. *EMBO J*. 15:6605-6616); the *plo1* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *byr3* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *byr4* gene product from *Saccharomyces cerevisiae* (Cullen, C. F., et al. 2000. A new genetic method for isolating functionally interacting genes: high plo1(+)-dependent mutants and their suppressors define genes in mitotic and septation pathways in fission yeast. *Genetics*. 155:1521-1534); the *pds1* gene product from *Saccharomyces cerevisiae* (Yamamoto, A., et al. 1996. Pds1p, an inhibitor of anaphase in budding yeast, plays a critical role in the APC and checkpoint pathway(s). *J. Cell Biol.* 133:99-110); the *esp1* gene product from *Saccharomyces cerevisiae* (Rao, H., et al. 2001. Degradation of a cohesin subunit by the N-end rule pathway is essential for chromosome stability. *Nature*. 410:955-999); the *ycs4* gene product from *Saccharomyces cerevisiae* (Biggins, S., et al. 2001. Genes involved in sister chromatid separation and segregation in the budding yeast *Saccharomyces cerevisiae*. *Genetics*. 159:453-470); the *cse4* gene product from *Saccharomyces cerevisiae* (Stoler, S. et al. 1995. A mutation in CSE4, an essential gene encoding a novel chromatin-associated protein in yeast, causes chromosome nondisjunction and cell cycle arrest at mitosis. *Genes Dev.* 9:573-586); the *ip11* gene product from *Saccharomyces cerevisiae* (Biggins, S., and A. W. Murray. 2001. The budding yeast protein kinase Ip11/Aurora allows the absence of tension to activate the spindle checkpoint. *Genes Dev.* 15:3118-3129); the *smt3* gene product from *Saccharomyces cerevisiae* (Takahashi, Y., et al. 1999. Smt3, a SUMO-1 homolog, is conjugated to Cdc3, a component of septin rings at the mother-bud neck in budding yeast. *Biochem. Biophys. Res. Commun.* 259:582-587); the *prp16* gene product from *Saccharomyces cerevisiae* (Hotz, H. R., and B. Schwer. 1998. Mutational analysis of the yeast DEAH-box splicing factor Prp16. *Genetics*. 149:807-815); the *prp19* gene product from *Saccharomyces cerevisiae* (Chen, C. H., et al. 2001. Identification and characterization of two novel components of the Prp19p-associated complex, Ntc30p and Ntc20p. *J. Biol. Chem.* 276:488-494); the *wss1* gene product from *Saccharomyces cerevisiae* (Biggins, S., et al. 2001. Genes involved in sister chromatid separation and segregation in the budding yeast *Saccharomyces cerevisiae*. *Genetics*. 159:453-470); the histone H4 gene product from *Saccharomyces cerevisiae* (Smith, M. M., et al. 1996. A novel histone H4 mutant defective in nuclear division and mitotic chromosome transmission. *Mol. Cell Biol.* 16:1017-1026); the histone H3 gene product from *Saccharomyces cerevisiae* (Smith, M. M., et al. 1996. A novel histone H14 mutant defective in nuclear division and mitotic chromosome transmission. *Mol. Cell Biol.* 16:1017-1026); the *cse4* gene product from *Saccharomyces cerevisiae* (Stoler, S., et al. 1995. A mutation in CSE4, an essential gene encoding a novel chromatin-associated protein in yeast, causes chromosome nondisjunction and cell cycle arrest at mitosis. *Genes Dev.* 9:573-586); the *spt4* gene product from *Saccharomyces cerevisiae* (Basrai, M. A., et al. 1996. Faithful chromosome transmission requires Spt4p, a putative regulator of chromatin structure in *Saccharomyces cerevisiae*. *Mol. Cell Biol.* 16:2838-2847); the *spt5* gene product from *Saccharomyces cerevisiae* (Yamaguchi, Y., et al. 2001. SPT genes: key players in the regulation of transcription, chromatin structure and other cellular processes. *J. Biochem. (Tokyo)*. 129:185-191); the *spt6* gene product from *Saccharomyces cerevisiae* (Clark-Adams, C. D., and F. Winston. 1987. The SPT6 gene is essential for growth and is required for delta-mediated transcription in *Saccharomyces cerevisiae*. *Mol. Cell Biol.* 7:679-686); the *ndc10* gene product from *Saccharomyces cerevisiae* (Chiang, P. W., et al. 1998. Isolation of murine SPT5 homologue: completion of the isolation and characterization of human and murine homologues of yeast chromatin structural protein complex SPT4, SPT5, and SPT6. *Genomics*. 47:426-428); the *ctf13* gene product from *Saccharomyces cerevisiae* (Doheny et al., Identification of essential components of the *S. cerevisiae* kinetochore, *Cell*

73:761-774, 1993); the spo1 gene product from *Saccharomyces cerevisiae* (Tavormina et al. 1997. Differential requirements for DNA replication in the activation of mitotic checkpoints in *Saccharomyces cerevisiae*. Mol. Cell Biol. 17:3315-3322); the cwp1 gene product from *Saccharomyces cerevisiae* (Tevzadze, G. G., et al. 2000. Spo1, a phospholipase B homolog, is required for spindle pole body duplication during meiosis in *Saccharomyces cerevisiae*. Chromosoma. 109:72-85); the dhp1 gene product from *Schizosaccharomyces pombe* (Shobuike, T., et al. 2001. The dhp1(+) gene, encoding a putative nuclear 5'→3' exoribonuclease, is required for proper chromosome segregation in fission yeast. Nucleic Acids Res. 29:1326-1333); the rat1 gene product from *Saccharomyces cerevisiae* (Shobuike, T., et al. 2001. The dhp1(+) gene, encoding a putative nuclear 5'→3' exoribonuclease, is required for proper chromosome segregation in fission yeast. Nucleic Acids Res. 29:1326-1333); the hsk1 gene product from *Saccharomyces cerevisiae* (Masai, H., et al. 1995. hsk1+, a *Schizosaccharomyces pombe* gene related to *Saccharomyces cerevisiae* CDC7, is required for chromosomal replication. EMBO J. 14:3094-3104); the dfp1 gene product from *Saccharomyces cerevisiae* (Takeda, T., et al. 1999. A fission yeast gene, him1(+)/dfp1(+), encoding a regulatory subunit for Hsk1 kinase, plays essential roles in S-phase initiation as well as in S-phase checkpoint control and recovery from DNA damage. Mol. Cell Biol. 19:5535-5547); the dbf4 gene product from *Saccharomyces cerevisiae* (Weinreich, M., and B. Stillman. 1999. Cdc7p-Dbf4p kinase binds to chromatin during S phase and is regulated by both the APC and the RAD53 checkpoint pathway. EMBO J. 18:5334-5346); the rad53 gene product from *Saccharomyces cerevisiae* (Sun, Z., et al. Spk1/Rad53 is regulated by Mec1-dependent protein phosphorylation in DNA replication and damage checkpoint pathways. Genes Dev. 10:395-406); the ibd1 gene product from *Saccharomyces cerevisiae* (Lee, J., et al. 1999. Ibd1p, a possible spindle pole body associated protein, regulates nuclear division and bud separation in *Saccharomyces cerevisiae*. Biochim. Biophys. Acta. 1449:239-253); and the hrp1 gene product from *Saccharomyces cerevisiae* (Henry, M., et al. 1996. Potential RNA binding proteins in *Saccharomyces cerevisiae* identified as suppressors of temperature-sensitive mutations in NPL3. Genetics. 142:103-115).

[0307] II.B. Gene Expression in Minicells

[0308] II.B.1. In General

[0309] In some aspects of the invention, it may be desirable to alter the expression of a gene and the production of the corresponding gene product. As is known in the art, and is used herein, a "gene product" may be a protein (polypeptide) or nucleic acid. Gene products that are proteins include without limitation enzymes, receptors, transcription factors, termination factors, expression factors, DNA-binding proteins, proteins that effect nucleic acid structure, or subunits of any of the preceding. Gene products that are nucleic acids include, but are not limited to, ribosomal RNAs (rRNAs), transfer RNAs (tRNAs), antisense RNAs, nucleases (including but not limited to catalytic RNAs, ribonucleases, and the like).

[0310] Depending on the function of a gene product, and on the type of application of the invention, it may be desirable to increase protein production, decrease protein

production, increase protein nucleic acid production and/or increase nucleic acid production. Provided herein are non-limiting examples of genes and gene products that may be manipulated, individually or in combination, in order to modulate the expression of gene products to be included into minicells or parent strains from which minicells are derived. The expression elements so modulated may be chromosomal and/or episomal, and may be expressed constitutively or in a regulated fashion, i.e., repressible and/or inducible. Furthermore, gene products under the regulation may be either monocistronic or polycistronic with other genes or with themselves.

[0311] II.B.2. Protein Production

[0312] By way of non-limiting example, increased protein production may occur through increased gene dosage (increased copy number of a given gene under the control of the native or artificial promoter where this gene may be on a plasmid or in more than one copy on the chromosome), modification of the native regulatory elements, including, but not limited to the promoter or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/enhancers, or relevant antisense nucleic acid or nucleic acid analog, cloning on a plasmid under the control of the native or artificial promoter, and increased or decreased production of native or artificial promoter regulatory element(s) controlling production of the gene or gene product.

[0313] By way of non-limiting example, decreased protein production may occur through modification of the native regulatory elements, including, but not limited to the promoter or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/enhancers, or relevant antisense nucleic acid or nucleic acid analog, through cloning on a plasmid under the control of the native regulatory region containing mutations or an artificial promoter, either or both of which resulting in decreased protein production, and through increased or decreased production of native or artificial promoter regulatory element(s) controlling production of the gene or gene product.

[0314] As used herein with regards to proteins, "intramolecular activity" refers to the enzymatic function or structure-dependent function. By way of non-limiting example, alteration of intramolecular activity may be accomplished by mutation of the gene, in vivo or in vitro chemical modification of the protein, inhibitor molecules against the function of the protein, e.g. competitive, non-competitive, or uncompetitive enzymatic inhibitors, inhibitors that prevent protein-protein, protein-nucleic acid, or protein-lipid interactions, e.g. expression or introduction of dominant-negative or dominant-positive protein or other protein fragment(s), carbohydrate(s), fatty acid(s), lipid(s), and nucleic acid(s) that may act directly or allosterically upon the protein, and/or modification of protein, carbohydrate, fatty acid, lipid, or nucleic acid moieties that modify the gene or gene product to create the functional protein.

[0315] As used herein with regards to proteins, "intermolecular function" refers to the effects resulting from an intermolecular interaction between the protein or nucleic acid and another protein, carbohydrate, fatty acid, lipid, nucleic acid, or other molecule(s) in or on the cell or the action of a product or products resulting from such an



interaction. By way of non-limiting example, intermolecular or intramolecular function may be the act or result of intermolecular phosphorylation, biotinylation, methylation, acylation, glycosylation, and/or other signaling event; this function may be the result of a protein-protein, protein-nucleic acid, or protein-lipid complex, and/or carrier function, e.g. the capacity to bind, either covalently or non-covalently small organic or inorganic molecules, protein(s), carbohydrate(s), fatty acid(s), lipid(s), and nucleic acid(s); this function may be to interact with the membrane to recruit other molecules to this compartment of the cell; this function may be to regulate the transcription and/or translation of the gene, other protein, or nucleic acid; and this function may be to stimulate the function of another process that is not yet described or understood.

### [0316] II.B.3. Nucleic Acid Production

[0317] By way of non-limiting example, increased nucleic acid production may occur through increased gene dosage (increased copy number of a given gene under the control of the native or artificial promoter where this gene may be on a plasmid or in more than one copy on the chromosome), modification of the native regulatory elements, including, but not limited to the promoter or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/enhancers, or relevant antisense nucleic acid or nucleic acid analog, cloning on a plasmid under the control of the native or artificial promoter, and increased or decreased production of native or artificial promoter regulatory element(s) controlling production of the gene or gene product.

[0318] By way of non-limiting example, decreased nucleic acid production may occur through modification of the native regulatory elements, including, but not limited to the promoter or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/enhancers, or relevant antisense nucleic acid or nucleic acid analog, through cloning on a plasmid under the control of the native regulatory region containing mutations or an artificial promoter, either or both of which resulting in decreased protein production, and through increased or decreased production of native or artificial promoter regulatory element(s) controlling production of the gene or gene product.

[0319] As used herein with regards to nucleic acids, "intramolecular activity" refers to a structure-dependent function. By way of non-limiting example, alteration of intramolecular activity may be accomplished by mutation of the gene, in vivo or in vitro chemical modification of the nucleic acid, inhibitor molecules against the function of the nucleic acid, e.g. competitive, non-competitive, or uncompetitive enzymatic inhibitors, inhibitors that prevent protein-nucleic acid interactions, e.g. expression or introduction of dominant-negative or dominant-positive protein or other nucleic acid fragment(s), or other carbohydrate(s), fatty acid(s), and lipid(s) that may act directly or allosterically upon the nucleic acid or nucleic acid-protein complex, and/or modification of nucleic acid moieties that modify the gene or gene product to create the functional nucleic acid.

[0320] As used herein with regards to nucleic acids, "intermolecular function" refers to the effects resulting from an intermolecular interaction between the nucleic acid and another nucleic acid, protein, carbohydrate, fatty acid, lipid,

or other molecule(s) in or on the cell or the action of a product or products resulting from such an interaction. By way of non-limiting example, intermolecular function may be the act or result of intermolecular or intramolecular phosphorylation, biotinylation, methylation, acylation, glycosylation, and/or other signaling event; this function may be the result of a protein-nucleic acid, and/or carrier function, e.g. the capacity to bind, either covalently or non-covalently small organic or inorganic molecules, protein(s), carbohydrate(s), fatty acid(s), lipid(s), and other nucleic acid(s); this function may be to interact with the membrane to recruit other molecules to this compartment of the cell; this function may be to regulate the transcription and/or translation of the gene, other nucleic acid, or protein; and this function may be to stimulate the function of another process that is not yet described or understood.

### [0321] II.C. Genes and Gene Products for Regulation of Expression

[0322] As is known in the art, a variety of genes, gene products and expression elements may be manipulated, individually or in combination, in order to modulate the expression of genes and/or production gene products. These include, by way of non-limiting example, RNA polymerases, ribosomes (ribosomal proteins and ribosomal RNAs), transfer RNAs (tRNAs), amino transferases, regulatory elements and promoter regions, transportation of inducible and inhibitory compounds, catabolite repression, general deletions and modifications, cytoplasmic redox state, transcriptional terminators, mechanisms for ribosomal targeting, proteases, chaperones, export apparatus and membrane targeting, and mechanisms for increasing stability and solubility. Each of these is discussed in more detail in the following sections. II.C.1. RNA Polymerases

[0323] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include modification of an endogenous and/or introduction of an exogenous RNA polymerase. A *rpo* gene, or any other gene that encodes a RNA polymerase subunit product from *E. coli*, or homologs of this gene or its gene product found in other prokaryotes, eukaryotes, archaeobacteria or organelles (mitochondria, chloroplasts, plastids and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells.

[0324] The production or activity of a desired gene product may be increased by increasing the level and/or activity of an RNA polymerase that transcribes the gene product's cognate gene. The production or activity of a desired protein gene product may be increased by decreasing the level and/or activity of an RNA polymerase that transcribes a gene product that inhibits the production or function of the desired gene product.

[0325] As one example, manipulation of the *rpoA* (*phs*, *sez*) gene or gene product from *E. coli*, or homologs of this gene or gene product found in other members of the Prokaryotes, Eukaryotes, Archaeobacteria and/or organelles (e.g., mitochondria, chloroplasts, plastids and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells. In addition to



rpoA, *E. coli*. genes that encode RNA polymerase subunits include rpoB (ftsR, groN, nitB, rif, ron, stl, stv, tabD, sdgB, mbrD), rpoC (tabD), rpoD (alt), rpoE, rpoH (fam, hin, htpR), rpoN (glnF, ntrA), rpoS (abrD, dpeB, katF, nur), and rpoZ (spoS). See Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein; and Sanderson et al., "Linkage Map of *Salmonella typhimurium*, Edition VIII" Chapter 110 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1903-1999, and references cited therein.

[0326] Production of a desired gene product may be preferentially or selectively enhanced by the introduction of an exogenous RNA polymerase that specifically recognizes expression sequences that are operably linked to the corresponding gene. By way of non-limiting example, the use of a T7 RNA polymerase to selectively express genes present on expression elements that segregate into minicells is described herein.

#### [0327] II.C.2. Ribosomes

[0328] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include modification of endogenous, and/or addition of exogenous, ribosomes or ribosomal subunits. The techniques may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells.

[0329] As is known in the art, a ribosome includes both proteins (polypeptides) and RNA (rRNA). Thus, in the case of a gene that encodes a component of a ribosome, the gene product may be a protein or an RNA. For a review, see Noller et al., "Ribosomes," Chapter 13 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 1, pages 167-186, and references cited therein. For the sake of convenience, both ribosomal proteins and rRNAs are encompassed by the term "ribosomal subunits."

[0330] The production or activity of a desired protein gene product may be increased by increasing the level and/or activity of a ribosomal subunit that causes or enhances the translation of the desired protein. The production or activity of a desired protein gene product may be increased by decreasing the level and/or activity of a ribosomal subunit that causes or enhances translation of a protein that has a negative impact on the production or activity of the desired protein.

[0331] Exemplary ribosomal genes and gene products that may be manipulated include without limitation the *E. coli* genes rimB, rimC, rimD, rimE, rimF (res), rimG, rimH, rimI, rimJ (tcp), rimK, rimL; rplA, rplB, rplC, rplD, rplE, rplF, rplI, rplJ, rplK, rplL, rplM, rplN, rplO, rplP, rplQ, rplR, rplS, rplT, rplU, rplV, rplW, rplX, rplY, rpsA, rpsB, rpsC,

rpsE (eps, spc, spcA), rpsF (sdgH), rpsG, rpsH, rpsI, rpsJ (nusE), rpsK, rpsL (strA), rpsM, rpsN, rpsO, rpsP, rpsQ, rpsR, rpsS, rpsT, rpsU, rpsV; rrfA, rrfB, rrfC, rrfD, rrfE, rrfF (rrfDbeta, rrvD), rrfG, rrfH; rrlA, rrlB, rrlC, rrlD, rrlE, rrlG, rrlH; rrnA, rrnB (csqE, rrnB1), rrnC (cqsB), rrnD (cqsD), rrnE (rrnD1), rrnG, rrnH; rrsA, rrsB, rrsC, rrsD, rrsE, rrsG, rrsH, and their cognate gene products.

[0332] Homologs of ribosomal genes or gene products found in other members of the Prokaryotes, Eukaryotes, Archaeobacteria and organelles (including but not limited to mitochondria, chloroplasts, plastids, and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or segregated minicells. See, for example, Barkan, A. and M. Goldschmidt-Clermont, Participation of nuclear genes in chloroplast gene expression, (2000) *Biochimie* 82:559-572; Willhoeft, U., H. Bu, and E. Tannich, Analysis of cDNA Expressed sequence tags from *Entamoeba histolytica*: Identification of two highly abundant polyadenylated transcripts with no overt open reading frames, (March 1999) *Protist* 150:61-70; Emelyanov, V., Evolutionary relationship of Rickettsiae and mitochondria (February 2001) *FEBS Letters* 501:11-18; and Gray, M., G. Burger and B. Lang, Mitochondrial Evolution (March 1999) *Science* 283:1476-1481. Ribosomal RNA sequences from a multitude of organisms and organelles are available through the Ribosomal Database Project (Maidak et al., A new version of the RDP (Ribosomal Database Project) (1999) *Nucleic Acids Research* 27:171-173). An index of ribosomal proteins classified by families on the basis of sequence similarities is available on-line at <http://www.expasy.ch/cgi-bin/lists?ribosomp.txt>; see also (Ramakrishnan et al., Ribosomal protein structures: insights into the architecture, machinery and evolution of the ribosome, *TIBS* 23:208-212, 1998.

#### [0333] II.C.3. Transfer RNAs (tRNAs)

[0334] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include utilization and/or modification of endogenous and/or exogenous transfer RNAs (tRNAs). Manipulation of the tRNA genes or gene products from *E. coli*, or homologs of tRNA genes or gene products found in other members of the Prokaryotes, Eukaryotes, Archaeobacteria and organelles (including but not limited to mitochondria, chloroplasts, plastids, and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells.

[0335] Exemplary *E. coli* tRNA genes include, but are not limited to, the alaT (talA) gene, the alaU (talD) gene, the alaV gene, the alaW (alaW) gene, the alaX (alaW) gene, the argQ (alaV) gene, the argU (dnaY, pin) gene, the alaU (talD) gene, the argV (argV2) gene, the argW gene, the argX gene, the argY (argV) gene, the argZ (argV) gene, the asnT gene, the asnU gene, the asnV gene, the aspT gene, the aspU gene, the cystT gene, the glnU (supB) gene, the glnV (supE) gene, the glnW (supB) gene, the gltT (tgtB) gene, the gltU (tgtC) gene, the gltV (tgtE) gene, the gltW gene, the glyT (sumA) gene, the glyU (sufD, sumA, sumB, supT) gene, the glyV (ins, mutA) gene, the glyW (ins, mutC) gene, the glyX gene, the glyY gene, the his R (hisT) gene, the ileT gene, the ileU gene, the ileV gene, the ileX gene, the leuP (leuV) gene, the

leuQ (leuV) gene, the leuQ (leuV) gene, the leuT gene, the leuU gene, the leuV (leuV) gene, the leuW (feeB) gene, the leuX (supP) gene, the leuZ gene, the lysT gene, the lysV (supN) gene, the lysW gene, the metI (metT) gene, the metU (metT) gene, the metV (metZ) gene, the metW gene, the metY gene, the pheU (pheR, pheW) gene, the pheV gene, the proK (proV) gene, the proL (prow) gene, the proM (proU) gene, the serT (divE) gene, the serU (ftsM, supD, supH) gene, the serV (supD) gene, the serW gene, the serX (serW) gene, the thrT gene, the thrU gene, the thrV gene, the thrW gene, the trpT (supU) gene, the tyrT (supC) gene, the tyrU (supM) gene, the atyV (tyrT, tyrT) gene, the valT gene, the valU (valU) gene, the valV (val) gene, the valW (val) gene, the valX gene, and the valX gene (Komine et al., Genomic Organization and Physical Mapping of the Transfer RNA Genes in *Escherichia coli* K12. J. Mol. Biol. 212:579-598, 1990; Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*, 2<sup>nd</sup> Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein; Sanderson et al., "Linkage Map of *Salmonella typhimurium*, Edition VIII" Chapter 110, Id., pages 1903-1999, and references cited therein; and Hershey, "Protein Synthesis," Chapter 40 in: *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*, Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1987, Volume 2, pages 613-647, and references cited therein).

[0336] Also included in the modification of transfer RNA molecules are the transfer RNA processing enzymes. Exemplary *E. coli* genes encoding tRNA processing enzymes include, but are not limited to the rnd gene (Blouin R T, Zaniewski R, Deutscher M P. Ribonuclease D is not essential for the normal growth of *Escherichia coli* or bacteriophage T4 or for the biosynthesis of a T4 suppressor tRNA, J. Biol. Chem. 258:1423-1426, 1983) and the rnpAB genes (Kirsebom L A, Baer M F, Altman S., Differential effects of mutations in the protein and RNA moieties of RNase P on the efficiency of suppression by various tRNA suppressors, J. Mol. Biol. 204:879-888, 1988).

[0337] Also included in the modification of transfer RNA molecules are modifications in endogenous tmRNAs and/or the introduction of exogenous tmRNAs to minicells and/or their parent cells. The tmRNA (a.k.a. 10S RNA) molecules have properties of tRNAs and mRNAs combined in a single molecule. Examples of tmRNAs are described in Zwieb et al. (Survey and Summary: Comparative Sequence Analysis of tmRNA, Nucl. Acids Res. 27:21063-2071, 1999).

#### [0338] II.C.4. Aminoacyl Synthetases

[0339] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include utilization and/or modification of endogenous and/or exogenous aminoacyl synthetases and proteins that effect their production and/or activity. Aminoacyl synthetases are involved in "charging" a tRNA molecule, i.e., attaching a tRNA to its cognate amino acid. (Martinis et al., Aminoacyl-tRNA Synthetases: General Features and Relationships. Chapter 58 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biol-*

*ogy*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 1, pages 887-901) and references cited therein; (Grunberg-Manago, Regulation of the Expression of Aminoacyl-tRNA Synthetases and Translation. Chapter 91 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 1, pages 1432-1457), and references cited therein; and (Hershey, "Protein Synthesis," Chapter 40 in: *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*, Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1987, Volume 1, pages 613-647), and references cited therein.

[0340] By way of non-limiting example, manipulation of the *aat* gene or gene product from *E. coli*, or homologs of this gene or gene product found in other members of the Prokaryotes, Eukaryotes, Archaeobacteria and/or organelles (e.g., mitochondria, chloroplasts, plastids and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells (Bochner, B. R., and Savageau, M. A. 1979. Inhibition of growth by imidazole propionic acid: evidence in vivo for coordination of histidine catabolism with the catabolism of other amino acids. Mol. Gen. Genet. 168(1):87-95).

[0341] In addition to *aat*, other exemplary *E. coli* genes encoding aminoacyl synthetases include *alaS* (*act*, *ala-act*, *lovB*) (Buckel et al., Suppression of temperature-sensitive aminoacyl-tRNA synthetase mutations by ribosomal mutations: a possible mechanism. Mol. Gen. Genet. 149:51-61, 1976); *argS* (*lovB*) (Eriani et al., Isolation and characterization of the gene coding for *Escherichia coli* arginyl-tRNA synthetase. Nucleic Acids Res. 17:5725-36, 1989); *asnS* (*ics*, *tss*) (Yamamoto et al., Identification of a temperature-sensitive asparaginyl-transfer ribonucleic acid synthetase mutant of *Escherichia coli*. J. Bacteriol. 132:127-31, 1977); *aspS* (*tlS*) (Eriani et al., Aspartyl-tRNA synthetase from *Escherichia coli*: cloning and characterisation of the gene, homologs of its translated amino acid sequence with asparaginyl- and lysyl-tRNA synthetases. Nucleic Acids Res. 18:7109-18, 1990); *cysS* (Eriani et al., Cysteinylyl-tRNA synthetase: determination of the last *E. coli* aminoacyl-tRNA synthetase primary structure. Nucleic Acids Res. 19:265-9, 1991); *ginS* (Yamao et al., *Escherichia coli* glutaminylyl-tRNA synthetase. I. Isolation and DNA sequence of the *glnS* gene. J. Biol. Chem. 257:11639-43, 1982); *gltE* (Lapointe et al., Thermosensitive mutants of *Escherichia coli* K-12 altered in the catalytic Subunit and in a Regulatory factor of the glutamy-transfer ribonucleic acid synthetase. J. Bacteriol. 122:352-8, 1975); *gltM* (Lapointe et al., Thermosensitive mutants of *Escherichia coli* K-12 altered in the catalytic Subunit and in a Regulatory factor of the glutamy-transfer ribonucleic acid synthetase. J. Bacteriol. 122:352-8, 1975); *gltX* (Lapointe et al., Thermosensitive mutants of *Escherichia coli* K-12 altered in the catalytic Subunit and in a Regulatory factor of the glutamy-transfer ribonucleic acid synthetase. J. Bacteriol. 122:352-8, 1975); *glyQ* (*glySa*) (Webster et al., Primary structures of both subunits of *Escherichia coli* glycyl-tRNA synthetase, J. Biol. Chem. 252:10637-41, 1983); *glyS* (*act*, *gly*, *glySB*) (Id.); *his S* (Parker et al., Mapping his S, the structural gene for histidyl-transfer ribonucleic acid synthetase, in *Escherichia coli*. J.

Bacteriol. 138:264-7, 1979); ileS (Singer et al., Synthesis of the isoleucyl- and valyl-tRNA synthetases and the isoleucine-valine biosynthetic enzymes in a threonine deaminase regulatory mutant of *Escherichia coli* K-12. J. Mol. Biol. 175:39-55, 1984); leuS (Morgan et al., Regulation of biosynthesis of aminoacyl-transfer RNA synthetases and of transfer-RNA in *Escherichia coli*. Arch. Biol. Med. Exp. (Santiago.) 12:415-26, 1979); lysS (herC, asaD) (Clark et al., Roles of the two lysyl-tRNA synthetases of *Escherichia coli*: analysis of nucleotide sequences and mutant behavior. J. Bacteriol. 172:3237-43, 1990); lysU (Clark et al., Roles of the two lysyl-tRNA synthetases of *Escherichia coli*: analysis of nucleotide sequences and mutant behavior, J. Bacteriol. 172:3237-43, 1990); metG (Dardel et al., Molecular cloning and primary structure of the *Escherichia coli* methionyl-tRNA synthetase gene. J. Bacteriol. 160:1115-22, 1984); pheS (phe-act) (Elseviers et al., Molecular cloning and regulation of expression of the genes for initiation factor 3 and two aminoacyl-tRNA synthetases, J. Bacteriol. 152:357-62, 1982); pheT (Corner et al., Genes for the alpha and beta subunits of the phenylalanyl-transfer ribonucleic acid synthetase of *Escherichia coli*. J. Bacteriol. 127:923-33, 1976); proS (drp) (Bohman et al., A temperature-sensitive mutant in prolinyl-tRNA ligase of *Escherichia coli* K-12. Mo. Gen. Genet. 177:603-5, 1980); serS (Hartlein et al., Cloning and characterization of the gene for *Escherichia coli* seryl-tRNA synthetase. Nucleic Acids Res. 15:1005-17, 1987); thrS (Frohler et al., Genetic analysis of mutations causing borrelidin resistance by overproduction of threonyl-transfer ribonucleic acid synthetase. J. Bacteriol. 143:1135-41, 1980); trpS (Hall et al., Cloning and characterization of the gene for *Escherichia coli* tryptophanyl-transfer ribonucleic acid synthetase. J. Bacteriol. 148:941-9, 1981); tyrS (Buonocore et al., Properties of tyrosyl transfer ribonucleic acid synthetase from two tyrS mutants of *Escherichia coli* K-12. J. Biol. Chem. 247:4843-9, 1972); and valS (Baer et al., Regulation of the biosynthesis of aminoacyl-transfer ribonucleic acid synthetases and of transfer ribonucleic acid in *Escherichia coli*. V. Mutants with increased levels of valyl-transfer ribonucleic acid synthetase. J. Bacteriol. 139:165-75, 1979).

#### [0342] II.C.5. Regulatory Elements and Promoter Regions

[0343] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include utilization and/or modification of regulatory elements and promoter regions. Such manipulations may result in increased or decreased production, and/or changes in the intramolecular and intermolecular functions, of a protein in a segregated minicell or its parent cell prior to minicell formation; in the latter instance, the protein may be one that is desirably retained in segregated minicells.

[0344] The production or activity of a desired gene product may be increased by increasing the level and/or activity of a promoter or other regulatory region that acts to stimulate or enhance the production of the desired gene product. The production or activity of a desired gene product may be increased by decreasing the level and/or activity of a promoter or other regulatory region that acts to stimulate or enhance the production of a gene product that acts to reduce or eliminate the level and/or activity of the desired gene product.

#### [0345] II.C.5.a. *Escherichia coli*

[0346] Regulatory elements, promoters and other expression elements and expression factors from *E. coli* include but are not limited to acrR (Ma, D., et al. 1996. The local repressor AcrR plays a modulating role in the regulation of acrAB genes of *Escherichia coli* by global stress signals. Mol. Microbiol. 19:101-112); ampD (Lindquist, S., et al. 1989. Signalling proteins in enterobacterial AmpC beta-lactamase regulation. Mol. Microbiol. 3:1091-1102; Holtje, J. V., et al. 1994. The negative regulator of beta-lactamase induction AmpD is a N-acetyl-anhydromuramyl-L-alanine amidase. FEMS Microbiol. Lett. 122:159-164); appR (Diaz-Guerra, L., et al. 1989. appR gene product activates transcription of microcin C7 plasmid genes. J. Bacteriol. 171:2906-2908; Touati, E., et al. 1991. Are appR and katF the same *Escherichia coli* gene encoding a new sigma transcription initiation factor? Res. Microbiol. 142:29-36); appY (Atlung, T., et al. 1989. Isolation, characterization, and nucleotide sequence of appY, a regulatory gene for growth-phase-dependent gene expression in *Escherichia coli*. J. Bacteriol. 171:1683-1691); araC (Casadaban, M. J., et al. 1976. Regulation of the regulatory gene for the arabinose pathway, araC. J. Mol. Biol. 104:557-566); arcA (luchi, S., and E. C. Lin. 1988. arcA (dye), a global regulatory gene in *Escherichia coli* mediating repression of enzymes in aerobic pathways. Proc. Natl. Acad. Sci. 85:1888-1892; luchi, S., et al. 1989. Differentiation of arcA, arcB, and cpxA mutant phenotypes of *Escherichia coli* by sex pilus formation and enzyme regulation. J. Bacteriol. 171:2889-2893); argR (xerA, Rarg) (Kelln, R. A., and V. L. Zak. 1978. Arginine regulon control in a *Salmonella typhimurium*-*Escherichia coli* hybrid merodiploid. Mol. Gen. Genet. 161:333-335; Vogel, R. H., et al. 1978. Evidence for translational repression of arginine biosynthetic enzymes in *Escherichia coli*: altered regulation in a streptomycin-resistant mutant. Mol. Gen. Genet. 162:157-162); ascG (Hall, B. G., and L. Xu. Nucleotide sequence, function, activation, and evolution of the cryptic asc operon of *Escherichia coli* K12. Mol. Biol. Evol. 9:688-706); aslB (Bennik, M. H., et al. 2000. Defining a rob regulon in *Escherichia coli* by using transposon mutagenesis. J. Bacteriol. 182:3794-3801); asnC (Kolling, R., and H. Lother. 1985. AsnC: an autogenously regulated activator of asparagine synthetase A transcription in *Escherichia coli*. J. Bacteriol. 164:310-315); atoC (Jenkins, L. S., and W. D. Nunn. 1987. Regulation of the ato operon by the atoC gene in *Escherichia coli*. J. Bacteriol. 169:2096-2102); baeR (Nagasawa, S., et al. 1993. Novel members of the two-component signal transduction genes in *Escherichia coli*. J. Biochem. (Tokyo). 114:350-357); baeS (Id.Id.); barA (Nagasawa, S., et al. 1992. A novel sensor-regulator protein that belongs to the homologous family of signal-transduction proteins involved in adaptive responses in *Escherichia coli*. Mol. Microbiol. 6:799-807; Ishige, K., et al. 1994. A novel device of bacterial signal transducers. EMBO J. 13:5195-5202); basS (Nagasawa, S., et al. 1993. Novel members of the two-component signal transduction genes in *Escherichia coli*. J. Biochem. (Tokyo). 114:350-357); betI (Lamark, T., et al. 1996. The complex bet promoters of *Escherichia coli*: regulation by oxygen (ArcA), choline (BetI), and osmotic stress. J. Bacteriol. 178:1655-1662); bglG (bglC, bglS) (Schnetz, K., and B. Rak. 1988. Regulation of the bgl operon of *Escherichia coli* by transcriptional antitermination. EMBO J. 7:3271-3277; Schnetz, K., and B. Rak. 1990. Beta-glucoside permease represses the bgl

- operon of *Escherichia coli* by phosphorylation of the anti-terminator protein and also interacts with glucose-specific enzyme III, the key element in catabolite control. Proc. Natl. Acad. Sci. 87:5074-5078); birA (bioR, dhhB) (Barker, D. F., and A. M. Campbell. 1981. Genetic and biochemical characterization of the birA gene and its product: evidence for a direct role of biotin holoenzyme synthetase in repression of the biotin operon in *Escherichia coli*. J. Mol. Biol. 146:469-492; Barker, D. F., and A. M. Campbell. 1981. The birA gene of *Escherichia coli* encodes a biotin holoenzyme synthetase. J. Mol. Biol. 146:451-467; Howard, P. K., et al. 1985. Nucleotide sequence of the birA gene encoding the biotin operon repressor and biotin holoenzyme synthetase functions of *Escherichia coli*. Gene. 35:321-331); btuR (Lundrigan, M. D., et al. 1987. Separate regulatory systems for the repression of metE and btuB by vitamin B12 in *Escherichia coli*. Mol. Gen. Genet. 206:401-407; Lundrigan, M. D., and R. J. Kadner. 1989. Altered cobalamin metabolism in *Escherichia coli* btuR mutants affects btuB gene regulation. J. Bacteriol. 171:154-161); cadC (Watson, N., et al. 1992. Identification of elements involved in transcriptional regulation of the *Escherichia coli* cad operon by external pH. J. Bacteriol. 174:530-540); celD (Parker, L. L., and B. G. Hall. 1990. Characterization and nucleotide sequence of the cryptic cel operon of *Escherichia coli* K12. Genetics. 124:455-471); chaB (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli and Salmonella typhimurium: cellular and molecular biology*, 2nd ed. American Society for Microbiology, Washington D.C.); chaC (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. 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A trans-acting regulatory mutation that causes overproduction of phosphatidylserine synthase in *Escherichia coli*. J. Biol. Chem. 258:9963-9967); purR (Meng, L. M., et al. 1990. Autoregulation of PurR repressor synthesis and involvement of purR in the regulation of purB, purC, purL, purMN and guaBA expression in *Escherichia coli*. Eur. J. Biochem. 187:373-379); putA (poaA) gene product from *Salmonella enterica* serotype *Typhimurium* (Menzel, R., and J. Roth. 1981. Regulation of the genes for proline utilization in *Salmonella typhimurium*: autogenous repression by the putA gene product. J. Mol. Biol. 148:21-44); pyrI (Cunin, R., et al. 1985. Structure-function relationship in allosteric aspartate carbamoyltransferase from *Escherichia coli*. I. Primary structure of a pyrI gene encoding a modified regulatory subunit. J. Mol. Biol. 186:707-713); rbsR (Lopilato, J. E., et al. 1984. D-ribose metabolism in *Escherichia coli* K-12: genetics, regulation, and transport. J. Bacteriol. 158:665-673); rcsA (Gottesman, S., et al. 1985. Regulation of capsular polysaccharide synthesis in *Escherichia coli* K-12: characterization of three regulatory genes. J. Bacteriol. 162:1111-1119); rcsB (Id.); rcsC (Id.); rcsF (Grevais, F. G., and G. R. Drapeau. 1992. Identification, cloning, and characterization of rcsF, a new regulator gene for exopolysaccharide synthesis that suppresses the division mutation ftsZ84 in *Escherichia coli* K-12. J. Bacteriol. 174:8016-8022); relB (Christensen, S. K., et al. 2001. RelE, a global inhibitor of translation, is activated during nutritional stress. Proc Natl. Acad. Sci. 98:14328-14333); rfaH (sfrB) (Pradel, E., and C. A. Schnaitman. 1991. Effect of rfaH (sfrB) and temperature on expression of rfa genes of *Escherichia coli* K-12. J. Bacteriol. 173:6428-6431); rhaR (Tobin, J. F., and R. F. Schleif. 1987. 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Effects of the *Escherichia coli* sfsA gene on mal genes expression and a DNA binding activity of SfsA. Biosci. Biotechnol. Biochem. 65:213-217); sfsB (nlp, sfs1) (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli and Salmonella typhimurium: cellular and molecular biology*, 2nd ed. American Society for Microbiology, Washington D.C.); soxR (Tsaneva, I. R., and B. Weiss. 1990. soxR, a locus governing a superoxide response regulon in *Escherichia coli* K-12. J. Bacteriol. 172:4197-4205); soxS (Wu, J., and B. Weiss. 1991. Two divergently transcribed genes, soxR and soxS, control a superoxide response regulon of



- Escherichia coli*. J. Bacteriol. 173:2864-2871); srlR (gutR) (Csonka, L. N., and A. J. Clark. 1979. Deletions generated by the transposon Tn10 in the srl recA region of the *Escherichia coli* K-12 chromosome. Genetics. 93:321-343); tdcA (Ganduri, Y. L., et al. 1993. TdcA, a transcriptional activator of the tdcABC operon of *Escherichia coli*, is a member of the LysR family of proteins. Mol. Gen. Genet. 240:395-402); tdcR (Hagewood, B. T., et al. 1994. Functional analysis of the tdcABC promoter of *Escherichia coli*: roles of TdcA and TdcR. J. Bacteriol. 176:6241-6220); thrS (Springer, M., et al. 1985. Autogenous control of *Escherichia coli* threonyl-tRNA synthetase expression in vivo. J. Mol. Biol. 185:93-104); tor R (Simon, G., et al. 1994. The tor R gene of *Escherichia coli* encodes a response regulator protein involved in the expression of the trimethylamine N-oxide reductase genes. J. Bacteriol. 176:5601-5606); treR (Horlacher, R., and W. Boos. 1997. Characterization of TreR, the major regulator of the *Escherichia coli* trehalose system. J. Biol. Chem. 272:13026-13032); trpR (Qunsalus, R. P., and C. Yanofsky. 1980. Nucleotide sequence and expression of *Escherichia coli* trpR, the structural gene for the trp aporepressor. Proc. Natl. Acad. Sci. 77:7117-7121); tyrR (Camakaris, H., and J. Pittard. 1973. Regulation of tyrosine and phenylalanine biosynthesis in *Escherichia coli* K-12: properties of the tyrR gene product. J. Bacteriol. 115:1135-1144); uhpA (Kadner, R. J., and D. M. Shattuck-Eidens. 1983. Genetic control of the hexose phosphate transport system of *Escherichia coli*: mapping of deletion and insertion mutations in the uhp region. J. Bacteriol. 155:1052-1061); uidR (gusR) (Novel, M., and G. Novel. 1976. Regulation of beta-glucuronidase synthesis in *Escherichia coli* K-12: pleiotropic constitutive mutations affecting uxu and uidA expression. J. Bacteriol. 127:418-432); uspA (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli* and *Salmonella typhimurium: cellular and molecular biology*, 2nd ed. American Society for Microbiology, Washington D.C.); uxuR (Novel, M., and G. Novel. 1976. Regulation of beta-glucuronidase synthesis in *Escherichia coli* K-12: pleiotropic constitutive mutations affecting uxu and uidA expression. J. Bacteriol. 127:418-432); wrbA (Yang, W., et al. 1993. A stationary-phase protein of *Escherichia coli* that affects the mode of association between the trp repressor protein and operator-bearing DNA. Proc. Natl. Acad. Sci. 90:5796-5800); xapR (pndR) (Seeger, C., et al. 1995. Identification and characterization of genes (xapA, xapB, and xapR) involved in xanthosine catabolism in *Escherichia coli*. J. Bacteriol. 177:5506-5516); and xylR (Inouye, S., et al. 1987. Expression of the regulatory gene xylS on the TOL plasmid is positively controlled by the xylR gene product. Proc. Natl. Acad. Sci. 84:5182-5186);
- [0347] Regulatory elements, promoters and other expression elements and factors from prokaryotes other than *E. coli* and *B. subtilis* include without limitation ahyRI gene product from *Aeromonas hydrophila* and *Aeromonas salmonicida* (Swift, S., et al. 1997. Quorum sensing in *Aeromonas hydrophila* and *Aeromonas salmonicida*: identification of the LuxRI homologs AhyRI and AsaRI and their cognate N-acylhomoserine lactone signal molecules. J. Bacteriol. 179:5271-5281); angR gene product from *Vibrio anguillarum* (Salinas, P. C., et al. 1989. Regulation of the iron uptake system in *Vibrio anguillarum*: evidence for a cooperative effect between two transcriptional activators. Proc. Natl. Acad. Sci. 86:3529-3522); aphA gene product from *Vibrio cholerae* (Kovachikova, G., and K. Skorupski. 2001. Overlapping binding sites for the virulence gene regulators AphA, AphB and cAMP-CRP at the *Vibrio cholerae* tcpPH promoter. Mol. Microbiol. 41:393-407); aphB gene product from *Vibrio cholerae* (Kovachikova, G., and K. Skorupski. 2000. Differential activation of the tcpPH promoter by AphB determines biotype specificity of virulence gene expression in *Vibrio cholerae*. J. Bacteriol. 182:3228-3238); comE gene product from *Streptococcus pneumoniae* (Ween, O., et al. 1999. Identification of DNA binding sites for ComE, a key regulator of natural competence in *Streptococcus pneumoniae*. Mol. Microbiol. 33:817-827); esaI gene product from *Pantoea stewartii* subsp. *stewartii* (von Bodman, S. B., et al. 1998. A negative regulator mediates quorum-sensing control of exopolysaccharide production in *Pantoea stewartii* subsp. *stewartii*. Proc. Natl. Acad. Sci. 95:7687-7692); esaR gene product from *Pantoea stewartii* subsp. *stewartii* (Id.); expi gene product from *Erwinia chrysanthemi* (Nasser, W., et al. 1998. Characterization of the *Erwinia chrysanthemi* expl-expR locus directing the synthesis of two N-acyl-homoserine lactone signal molecules. Mol. Microbiol. 29:1391-1405); expR gene product from *Erwinia chrysanthemi* (Id.); gacA gene product from *Pseudomonas aeruginosa* (Pessi, G., and D. Haas. 2001. Dual control of hydrogen cyanide biosynthesis by the global activator GacA in *Pseudomonas aeruginosa* PAO1. FEMS Microbiol. Lett. 200:73-78); hapR gene product from *Vibrio cholerae* (Jobling, M. G., and R. K. Holmes. Characterization of hapR, a positive regulator of the *Vibrio cholerae* HA/protease gene hap, and its identification as a functional homologue of the *Vibrio harveyi* luxR gene. Mol. Microbiol. 26:1023-1034); hlyR gene product from *Vibrio cholerae* (von Mechow, S., et al. 1985. Mapping of a gene that regulates hemolysin production in *Vibrio cholerae*. J. Bacteriol. 163:799-802); hupR gene product from *Vibrio vulnificus* (Litwin, C. M., and J. Quackenbush. 2001. Characterization of a *Vibrio vulnificus* LysR homologue, HupR, which regulates expression of the haem uptake outer membrane protein, HupA. Microb. Pathog. 31:295-307); lasR gene product from *Pseudomonas aeruginosa* (Gambella, M. J., and B. H. Iglewski. 1991. Cloning and characterization of the *Pseudomonas aeruginosa* lasR gene, a transcriptional activator of elastase expression. J. Bacteriol. 173:3000-3009); leuO gene product from *Salmonella enterica* serovar *Typhimurium* (Fang, M., and H. Y. Wu. 1998. A promoter relay mechanism for sequential gene activation. J. Bacteriol. 180:626-633); luxI gene product from *Vibrio cholerae* (Engebrecht, J., and M. Silverman. Nucleotide sequence of the regulatory locus controlling expression of bacterial genes for bioluminescence. Nucleic Acids Res. 15:10455-10467); luxO gene product from *Vibrio cholerae* (Bassler, B. L., et al. 1994. Sequence and function of LuxO, a negative regulator of luminescence in *Vibrio harveyi*. Mol. Microbiol. 12:403-412); luxR gene product from *Vibrio cholerae* (Engebrecht, J., and M. Silverman. Nucleotide sequence of the regulatory locus controlling expression of bacterial genes for bioluminescence. Nucleic Acids Res. 15:10455-10467); phzR gene product from *Pseudomonas aureofaciens* (Pierson, L. S., et al. 1994. Phenazine antibiotic biosynthesis in *Pseudomonas aureofaciens* 30-84 is regulated by PhzR in response to cell density. J. Bacteriol. 176:3966-3974); rhlR



gene product from *Pseudomonas aeruginosa* (Ochsner, U. A. et al. 1994. Isolation and characterization of a regulatory gene affecting rhamnolipid biosurfactant synthesis in *Pseudomonas aeruginosa*. J. Bacteriol. 176:2044-2054); rsmA gene product from *Erwinia carotovora* subsp. *carotovora* (Cui, Y., et al. 1995. Identification of a global repressor gene, rsmA, of *Erwinia carotovora* subsp. *carotovora* that controls extracellular enzymes, N-(3-oxohexanoyl)-L-homoserine lactone, and pathogenicity in soft-rotting *Erwinia* spp. J. Bacteriol. 177:5108-5115); rsmB gene product from *Erwinia carotovora* subsp. *carotovora* (Cui, Y., et al. 1999. rsmC of the soft-rotting bacterium *Erwinia carotovora* subsp. *carotovora* negatively controls extracellular enzyme and harpin (Ecc) production and virulence by modulating levels of regulatory RNA (rsmB) and RNA-binding protein (RsmA). J. Bacteriol. 181:6042-6052); sirA gene product from *Salmonella enterica* serovar *Typhimurium* (Goodier, R. I., and B. M. Ahmer. 2001. SirA orthologs affects both motility and virulence. J. Bacteriol. 183:2249-2258); taf gene product from *Vibrio cholerae* (Salinas, P. C., et al. 1989. Regulation of the iron uptake system in *Vibrio anguillarum*: evidence for a cooperative effect between two transcriptional activators. Proc. Natl. Acad. Sci. 86:3529-3522); tcpP gene product from *Vibrio cholerae* (Hase, C. C., and J. J. Mekalanos. 1998. TcpP protein is a positive regulator of virulence gene expression in *Vibrio cholerae*. Proc. Natl. Acad. Sci. 95:730-734); toxR gene product from *Vibrio cholerae* (Miller, V. L., and J. J. Mekalanos. 1984. Synthesis of cholera toxin is positively regulated at the transcriptional level by toxR. Proc. Natl. Acad. Sci. 81:3471-4375); toxS gene product from *Vibrio cholerae* (Miller, V. L., et al. 1989. Identification of toxS, a regulatory gene whose product enhances toxR-mediated activation of the cholera toxin promoter. J. Bacteriol. 171:1288-1293); toxT from *Vibrio cholerae* (Kaufman, M. R., et al. 1993. Biogenesis and regulation of the *Vibrio cholerae* toxin-coregulated pilus: analogies to other virulence factor secretory systems. Gene. 126:43-49); traM gene product from *Agrobacterium tumefaciens* (Faqua, C., et al. 1995. Activity of the *Agrobacterium* Ti plasmid conjugal transfer regulator TraR is inhibited by the product of the traM gene. J. Bacteriol. 177:1367-1373); traR gene product from *Agrobacterium tumefaciens* (Piper, K. R., et al. 1993. Conjugation factor of *Agrobacterium tumefaciens* regulates Ti plasmid transfer by autoinduction. Nature. 362:448-450); vicH gene product from *Vibrio cholerae* (Tendeng, C., et al. 2000. Isolation and characterization of vicH, encoding a new pleiotropic regulator in *Vibrio cholerae*. J. Bacteriol. 182:2026-2032); vspR gene product from *Vibrio cholerae* (Yildiz, F. H., et al. 2001. VpsR, a Member of the Response Regulators of the Two-Component Regulatory Systems, Is Required for Expression of vps Biosynthesis Genes and EPS (ETr)-Associated Phenotypes in *Vibrio cholerae* O1 E1 Tor. J. Bacteriol. 183:1716-1726).

#### [0348] II.C.5. b. *Bacillus subtilis*

[0349] Regulatory elements, promoters and other expression elements and expression elements from *B. subtilis* include but are not limited to abrB (Perego, M., et al. 1988. Structure of the gene for the transition state regulator, abrB: regulator synthesis is controlled by the spo0A sporulation gene in *Bacillus subtilis*. Mol. Microbiol. 2:698-699); acoR (Ali, N. O., et al. 2001. Regulation of the acetoin catabolic pathway is controlled by sigma L in *Bacillus subtilis*. J. Bacteriol. 183:2497-2504); ahrC (Klinger, U., et al. 1995. A

binding site for activation by the *Bacillus subtilis* AhrC protein, a repressor/activator of arginine metabolism. Mol. Gen. Genet. 248:329-340); alaR (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); alsR (Renna, M. C., et al. 1993. Regulation of the *Bacillus subtilis* alsS, alsD, and alsR genes involved in post-exponential-phase production of acetoin. J. Bacteriol. 175:3863-3875); ansR (Sun, D., and P. Setlow. 1993. Cloning and nucleotide sequence of the *Bacillus subtilis* ansR gene, which encodes a repressor of the ans operon coding for L-asparaginase and L-aspartase. J. Bacteriol. 175:2501-2506); araR (Sa-Nogueira, I., and L. J. Mota. 1997. Negative regulation of L-arabinose metabolism in *Bacillus subtilis*: characterization of the araR (araC) gene. J. Bacteriol. 179:1598-1608); arfM (Marino, M., et al. 2001. Modulation of anaerobic energy metabolism of *Bacillus subtilis* by arfM (ywiD). J. Bacteriol. 183:6815-6821); arsR (Rosenstein, R., et al. 1992. Expression and regulation of the antimonite, arsenite, and arsenate resistance operon of *Staphylococcus xylosus* plasmid pSX267. J. Bacteriol. 174:3676-3683); azlB (Belitsky, B. R., et al. 1997. An Irp-like gene of *Bacillus subtilis* involved in branched-chain amino acid transport. J. Bacteriol. 179:54485457); birA (Bower, S., et al. 1995. Cloning and characterization of the *Bacillus subtilis* birA gene encoding a repressor of the biotin operon. J. Bacteriol. 177:2572-2575); bkdR (Bebarbouille, M., et al. 1999. Role of bkdR, a transcriptional activator of the sigL-dependent isoleucine and valine degradation pathway in *Bacillus subtilis*. J. Bacteriol. 181:2059-2066); bliR (Ahmed, M., et al. 1995. Two highly similar multidrug transporters of *Bacillus subtilis* whose expression is differentially regulated. J. Bacteriol. 177:3904-3910); bmrR (Ahmed, M., et al. 1994. A protein that activates expression of a multidrug efflux transporter upon binding the transporter substrates. J. Biol. Chem. 269:28506-28513); ccpA (Henkin, T. M., et al. 0.1991. Catabolite repression of alpha-amylase gene expression in *Bacillus subtilis* involves a trans-acting gene product homologous to the *Escherichia coli* lacI and galR repressors. Mol. Microbiol. 5:575-584); ccpB (Chauvaux, S., et al. 1998. CcpB, a novel transcription factor implicated in catabolite repression in *Bacillus subtilis*. J. Bacteriol. 180:491-497); ccpC (Jourlin-Castelli, C., et al. 2000. CcpC, a novel regulator of the LysR family required for glucose repression of the citB gene in *Bacillus subtilis*. J. Mol. Biol. 295:865-878); cggR (Fillinger, S., et al. 2000. Two glyceraldehyde-3-phosphate dehydrogenases with opposite physiological roles in a nonphotosynthetic bacterium. J. Biol. Chem. 275:14031-14037); cheB (Bischoff, D. S., and G. W. Ordal. 1991. Sequence and characterization of *Bacillus subtilis* CheB, a homolog of *Escherichia coli* CheY, and its role in a different mechanism of chemotaxis. J. Biol. Chem. 266:12301-12305); cheY (Bischoff, D. S., et al. 1993. Purification and characterization of *Bacillus subtilis* CheY. Biochemistry 32:9256-9261); citR (Jin, S., and A. L. Sohenshein. 1994. Transcriptional regulation of *Bacillus subtilis* citrate synthase genes. J. Bacteriol. 176:4680-4690); citT (Yamamoto, H., et al. 2000. The CitST two-component system regulates the expression of the Mg-citrate transporter in *Bacillus subtilis*. Mol. Microbiol. 37:898-912); codY (Slack, F. J., et al. 1995. A gene required for nutritional repression of the *Bacillus subtilis* dipeptide permease operon. Mol. Microbiol. 15:689-702); comA (Nakano, M. M., and P. Zuber. 1989. Cloning and characterization of

- srfB, a regulatory gene involved in surfactin production and competence in *Bacillus subtilis*. J. Bacteriol. 171:5347-5353); comK (Msadek, T., et al. 1994. MecB of *Bacillus subtilis*, a member of the ClpC ATPase family, is a pleiotropic regulator controlling competence gene expression and growth at high temperature. Proc. Natl. Acad. Sci. 91:5788-5792); comQ (Weinrauch, Y., et al. 1991. Sequence and properties of comQ, a new competence regulatory gene of *Bacillus subtilis*. J. Bacteriol. 173:5685-5693); cssR (Hyyrylainen, H. L., et al. 2001. A novel two-component regulatory system in *Bacillus subtilis* for the survival of severe secretion stress. Mol. Microbiol. 41:1159-1172); ctsR (Kruger, E., and M. Hecker. 1998. The first gene of the *Bacillus subtilis* clpC operon, ctsR, encodes a negative regulator of its own operon and other class III heat shock genes. J. Bacteriol. 180:6681-6688); dctR (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); degA (Bussey, L. B., and R. L. Switzer. 1993. The degA gene product accelerates degradation of *Bacillus subtilis* phosphoribosylpyrophosphate amidotransferase in *Escherichia coli*. J. Bacteriol. 175:6348-6353); degU (Msadek, T., et al. 1990. Signal transduction pathway controlling synthesis of a class of degradative enzymes in *Bacillus subtilis*: expression of the regulatory genes and analysis of mutations in degS and degU. J. Bacteriol. 172:824-834); deoR (Saxild, H. H., et al. 1996. Dra-nupC-pdp operon of *Bacillus subtilis*: nucleotide sequence, induction by deoxyribonucleosides, and transcriptional regulation by the deoR-encoded DeoR repressor protein. J. Bacteriol. 178:424-434); exuR (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); frn (Cruz Ramos, H., et al. 1995. Anaerobic transcription activation in *Bacillus subtilis*: identification of distinct FNR-dependent and -independent regulatory mechanisms. EMBO J. 14:5984-5994); fruR (Saier, M. H. Jr. 1996. Cyclic AMP-independent catabolite repression in bacteria. FEMS Microbiol. Lett. 138:97-103); fur (Chen, L., et al. 1993. Metalloregulation in *Bacillus subtilis*: isolation and characterization of two genes differentially repressed by metal ions. J. Bacteriol. 175:5428-5437); gabR (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); gerE (Holand, S. K., et al. 1987. The possible DNA-binding nature of the regulatory proteins, encoded by spoIID and gerE, involved in the sporulation of *Bacillus subtilis*. J. Gen. Microbiol. 133:2381-2391); glcR (Stulke, J., et al. 2001. 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- [0350] II.C.5.c. Other Eubacteria
- [0351] Regulatory elements, promoters and other expression elements and factors from prokaryotes other than *E. coli* and *B. subtilis* include without limitation ahyRI gene product from *Aeromonas hydrophila* and *Aeromonas salmoni-*

*cida* (Swift, S., et al. 1997. Quorum sensing in *Aeromonas hydrophila* and *Aeromonas salmonicida*: identification of the LuxRI homologs AhYRI and AsaRI and their cognate N-acylhomoserine lactone signal molecules. *J. Bacteriol.* 179:5271-5281); angR gene product from *Vibrio anguillarum* (Salinas, P. C., et al. 1989. Regulation of the iron uptake system in *Vibrio anguillarum*: evidence for a cooperative effect between two transcriptional activators. *Proc. Natl. Acad. Sci.* 86:3529-3522); aphA gene product from *Vibrio cholerae* (Kovachikova, G., and K. Skorupski. 2001. Overlapping binding sites for the virulence gene regulators AphA, AphB and cAMP-CRP at the *Vibrio cholerae* tcpPH promoter. *Mol. Microbiol.* 41:393-407); aphB gene product from *Vibrio cholerae* (Kovachikova, G., and K. Skorupski. 2000. Differential activation of the tcpPH promoter by AphB determines biotype specificity of virulence gene expression in *Vibrio cholerae*. *J. 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Bacteriol.* 177:5108-5115); rsmB gene product from *Erwinia carotovora* subsp. *carotovora* (Cui, Y., et al. 1999. rsmC of the soft-rotting bacterium *Erwinia carotovora* subsp. *carotovora* negatively controls extracellular enzyme and harpin(Ecc) production and virulence by modulating levels of regulatory RNA (rsmB) and RNA-binding protein (RsmA). *J. Bacteriol.* 181:6042-6052); sirA gene product from *Salmonella enterica* serovar *Typhimurium* (Goodier, R. I., and B. M. Ahmer. 2001. SirA orthologs affects both motility and virulence. *J. Bacteriol.* 183:2249-2258); taf gene product from *Vibrio cholerae* (Salinas, P. C., et al. 1989. Regulation of the iron uptake system in *Vibrio anguillarum*: evidence for a cooperative effect between two transcriptional activators. *Proc. Natl. Acad. Sci.* 86:3529-3522); tcpP gene product from *Vibrio cholerae* (Hase, C. C., and J. J. Mekalanos. 1998. TcpP protein is a positive regulator of virulence gene expression in *Vibrio cholerae*. *Proc. Natl. Acad. Sci.* 95:730-734); toxR gene product from *Vibrio cholerae* (Miller, V. L., and J. J. Mekalanos. 1984. Synthesis of cholera toxin is positively regulated at the transcriptional level by toxR. *Proc. Natl. Acad. Sci.* 81:3471-4375); toxS gene product from *Vibrio cholerae* (Miller, V. L., et al. 1989. Identification of toxS, a regulatory gene whose product enhances toxR-mediated activation of the cholera toxin promoter. *J. Bacteriol.* 171:1288-1293); toxT from *Vibrio cholerae* (Kaufman, M. R., et al. 1993. Biogenesis and regulation of the *Vibrio cholerae* toxin-coregulated pilus: analogies to other virulence factor secretory systems. *Gene.* 126:43-49); traM gene product from *Agrobacterium tumefaciens* (Faqua, C., et al. 1995. Activity of the *Agrobacterium* Ti plasmid conjugal transfer regulator TraR is inhibited by the product of the traM gene. *J. Bacteriol.* 177:1367-1373); traR gene product from *Agrobacterium tumefaciens* (Piper, K. R., et al. 1993. 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- solanacearum* (Van Gijsegem, F., et al. 1995. The hrp gene locus of *Pseudomonas solanacearum*, which controls the production of a type III secretion system, encodes eight proteins related to components of the bacterial flagellar biogenesis complex. *Mol. Microbiol.* 15:1095-1114); *carotovora* subsp. *carotovora* (Cui, Y., et al. 1995. Identification of a global repressor gene, rsmA, of *Erwinia carotovora* subsp. *carotovora* that controls extracellular enzymes, N-(3-oxohexanoyl)-L-homoserine lactone, and pathogenicity in soft-rotting *Erwinia* spp. *J. Bacteriol.* 177:5108-5115); rsmB gene product from *Erwinia carotovora* subsp. *carotovora* (Cui, Y., et al. 1999. rsmC of the soft-rotting bacterium *Erwinia carotovora* subsp. *carotovora* negatively controls extracellular enzyme and harpin(Ecc) production and virulence by modulating levels of regulatory RNA (rsmB) and RNA-binding protein (RsmA). *J. Bacteriol.* 181:6042-6052); sirA gene product from *Salmonella enterica serovar Typhimurium* (Goodier, R. I., and B. M. Ahmer. 2001. SirA orthologs affects both motility and virulence. *J. Bacteriol.* 183:2249-2258); taf gene product from *Vibrio cholerae* (Salinas, P. C., et al. 1989. Regulation of the iron uptake system in *Vibrio anguillarum*: evidence for a cooperative effect between two transcriptional activators. *Proc. Natl. Acad. Sci.* 86:3529-3522); tcpP gene product from *Vibrio cholerae* (Hase, C. C., and J. J. Mekalanos. 1998. TcpP protein is a positive regulator of virulence gene expression in *Vibrio cholerae*. *Proc. Natl. Acad. Sci.* 95:730-734); toxR gene product from *Vibrio cholerae* (Miller, V. L., and J. J. Mekalanos. 1984. Synthesis of cholera toxin is positively regulated at the transcriptional level by toxR. *Proc. Natl. Acad. Sci.* 81:3471-4375); toxS gene product from *Vibrio cholerae* (Miller, V. L., et al. 1989. Identification of toxS, a regulatory gene whose product enhances toxR-mediated activation of the cholera toxin promoter. *J. Bacteriol.* 171:1288-1293); toxT from *Vibrio cholerae* (Kaufman, M. R., et al. 1993. Biogenesis and regulation of the *Vibrio cholerae* toxin-coregulated pilus: analogies to other virulence factor secretory systems. *Gene.* 126:43-49); traM gene product from *Agrobacterium tumefaciens* (Faqua, C., et al. 1995. Activity of the *Agrobacterium* Ti plasmid conjugal transfer regulator TraR is inhibited by the product of the traM gene. *J. Bacteriol.* 177:1367-1373); traR gene product from *Agrobacterium tumefaciens* (Piper, K. R., et al. 1993. Conjugation factor of *Agrobacterium tumefaciens* regulates Ti plasmid transfer by autoinduction. *Nature.* 362:448-450); vicH gene product from *Vibrio cholerae* (Tendeng, C., et al. 2000. Isolation and characterization of vicH, encoding a new pleiotropic regulator in *Vibrio cholerae*. *J. Bacteriol.* 182:2026-2032); vspR gene product from *Vibrio cholerae* (Yildiz, F. H., et al. 2001. VpsR, a Member of the Response Regulators of the Two-Component Regulatory Systems, Is Required for Expression of vps Biosynthesis Genes and EPS(ETr)-Associated Phenotypes in *Vibrio cholerae* O1 E1 Tor. *J. Bacteriol.* 183:1716-1726); IrpA gene product from *Pyrococcus furiosus* (Brinkman, A. B., et al. 2000. An Lrp-like transcriptional regulator from the archaeon *Pyrococcus furiosus* is negatively autoregulated. *J. Biol. Chem.* 275:38160-38169); manR gene product from *Listeria monocytogenes* (Dalet, K., et al. 2001. A sigma(54)-dependent PTS permease of the mannose family is responsible for sensitivity of *Listeria monocytogenes* to mesentericin Y105. *Microbiology.* 147:3263-3269); msmR gene product from *Streptococcus mutans* (Russell, R. R., et al. 1992. A binding protein-dependent transport system in *Streptococcus mutans* responsible for multiple sugar metabolism. toxR gene product from *Vibrio cholerae* (Miller, V. L., and J. J. Mekalanos. 1984. Synthesis of cholera toxin is positively regulated at the transcriptional level by toxR. *Proc. Natl. Acad. Sci.* 81:3471-3475); padR gene product from *Pediococcus pentosaceus* (Barthelmebs, L., et al. 2000. Inducible metabolism of phenolic acids in *Pediococcus pentosaceus* is encoded by an autoregulated operon which involves a new class of negative transcriptional regulator. *J. Bacteriol.* 182:6724-6731); purR (Weng, M., et al. 1995); and xylR gene product from *Bacillus megaterium* (Rygus, T., et al. 1991. Molecular cloning, structure, promoters and regulatory elements for transcription of the *Bacillus megaterium* encoded regulon for xylose utilization. *Arch. Microbiol.* 155:535:542).
- [0352]** II.C.5.d. Bacteriophage and Transposable Elements
- [0353]** Regulatory elements, promoters and other expression elements from bacteriophage and transposable elements include without limitation cI gene product from bacteriophage lambda mation and/or segregated minicells (Reichardt, L. F. 1975. Control of bacteriophage lambda repressor synthesis: regulation of the maintenance pathway of the cro and cI products. *J. Mol. Biol.* 93:289-309); (Love, C. A., et al. 1996. Stable high-copy-number bacteriophage lambda promoter vectors for overproduction of proteins in *Escherichia coli*. *Gene.* 176:49-53); the c2 gene product from bacteriophage P22 (Gough, M., and S. Tokuno. 1975. Further structural and functional analogies between the repressor regions of phages P22 and lambda. *Mol. Gen. Genet.* 138:71-79); the cro gene from bacteriophage lambda (Reichardt, L. F. 1975. Control of bacteriophage lambda repressor synthesis: regulation of the maintenance pathway of the cro and cI products. *J. Mol. Biol.* 93:289-309); the ant gene from bacteriophage P22 (Youderian, P. et al. 1982. Sequence determinants of promoter activity. *Cell.* 30:843-853); the mnt gene from bacteriophage P22 (Gough, M. 1970. Requirement for a functional int product in temperature inductions of prophage P22 ts mnt. *J. Virol.* 6:320-325; Prell, H. H. 1978. Ant-mediated transactivation of early genes in *Salmonella* prophage P22 by superinfecting virulent P22 mutants. *Mol. Gen. Genet.* 164:331-334); the tetR gene product from the TetR family of bacterial regulators or homologues of this gene or gene product found in Tn10 and other members of the bacteriophage, animal virus, Eubacteria, Eucarya or Archaea may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or segregated minicells (Moyed, H. S., and K. P. Bertrand. 1983. Mutations in multicopy Tn10 tet plasmids that confer resistance to inhibitory effects of inducers of tet gene expression. *J. Bacteriol.* 155:557-564); the mnt gene product from bacteriophage SP6 mation and/or segregated minicells (Mead, D. A., et al. 1985. Single stranded DNA SP6 promoter plasmids for engineering mutant RNAs and proteins: synthesis of a 'stretched' preproparathyroid hormone. *Nucleic Acids Res.* 13:1103-1118); and the mnt gene product from bacteriophage T7 mation and/or segregated minicells (Steen, R., et al. 1986. T7 RNA polymerase directed expression of the *Escherichia coli* rrnB operon. *EMBO J.* 5:1099-1103).
- [0354]** II.C.5.e. Use of Site-Specific Recombination in Expression Systems
- [0355]** Included in the design of the invention are techniques that increase the efficiency of gene expression and

protein production in minicells. By way of non-limiting example, these techniques may include modification of endogenous and/or exogenous regulatory elements responsible for activation and/or repression of proteins to be expressed from chromosomal and/or plasmid expression vectors. By way of non-limiting example, this system may be applied to any of the above regulatory elements/systems. Specifically, each of the above mentioned regulatory systems may be constructed such that the promoter regions are oriented in a direction away from the gene to be expressed, or each of the above mentioned gene(s) to be expressed may be constructed such that the gene(s) to be expressed is oriented in a direction away from the regulatory region promoter. Constructed in this system is a methodology dependent upon site-specific genetic recombination for inversion and induction of the gene of interest (Backman, K., et al. 1984. Use of synchronous site-specific recombination in vivo to regulate gene expression. *Bio/Technology* 2:1045-1049; Balakrishnan, R., et al. 1994. A gene cassette for adapting *Escherichia coli* strains as hosts for att-Int-mediated rearrangement and pL expression vectors. *Gene* 138:101-104; Hasan, N., and W. Szybalaki. 1987. Control of cloned gene expression by promoter inversion in vivo: construction of improved vectors with a multiple cloning site and the Ptac promoter. *Gene* 56:145-151; Wulfig, C., and A. Pluckthun. 1993. A versatile and highly repressible *Escherichia coli* expression system based on invertible promoters: expression of a gene encoding a toxic gene product. *Gene* 136:199-203). These invertible promoters and/or gene regions will allow tight regulation of potentially toxic protein products. By way of non-limiting example, these systems may be derived from bacteriophage lambda, bacteriophage Mu, and/or bacteriophage P22. In any of these potential systems, regulation of the recombinase may be regulated by any of the regulatory systems discussed in section II.C.5 and elsewhere herein.

#### [0356] II° C.5.e. Use of Copy Number Control Switches

[0357] A method that can be used to increase the efficiency of gene expression and protein production in minicells involves the modification of endogenous and/or introduction of exogenous genetic expression systems such that the number of copies of a gene encoding a protein to be expressed can be modulated. Copy number control systems comprise elements designed to modulate copy number in a controlled fashion.

[0358] In an exemplary mode, copy number is controlled to decrease the effects of "leaky" (uninduced) expression of toxic gene products. This allows one to maintain the integrity of a potentially toxic gene product during processes such as cloning, culture maintenance, and periods of growth prior to minicell-induction. That is, decreasing the copy number of a gene is expected to decrease the opportunity for mutations affecting protein expression and/or function to arise. Immediately prior to, during and/or after minicell formation, the copy number may be increased to optimize the gene dosage in minicells as desired.

[0359] The replication of eubacterial plasmids is regulated by a number of factors, some of which are contained within the plasmid, others of which are located on the chromosome. For reviews, see del Solar, G., et al. 2000. Plasmid copy number control: an ever-growing story. *Mol Microbiol.* 37:492-500; del Solar, G., et al. 1998. Replication and

control of circular bacterial plasmids. *Microbiol Mol Biol Rev.* 62:434-64; and Filutowicz, M., et al. 1987. DNA and protein interactions in the regulation of plasmid replication. *J Cell Sci Suppl.* 7:15-31.

[0360] By way of non-limiting example, the *pcnB* gene product, the wildtype form of which promotes increased ColE1 plasmid copy number (Soderbom, F., et al. 1997. Regulation of plasmid R1 replication: *PcnB* and RNase E expedite the decay of the antisense RNA, *CopA*. *Mol. Microbiol.* 26:493-504), is modulated; and/or mutant forms of the *pcnB* gene are introduced into a cell. In an exemplary cell type that may be used in the methods of the invention, the wildtype *pcnB* chromosomal gene is replaced with a mutant *pcnB80* allele (Lopilato, J., et al. 1986. Mutations in a new chromosomal gene of *Escherichia coli* K-12, *pcnB*, reduce plasmid copy number of pBR322 and its derivatives. *Mol. Gen. Genet.* 205:285-290). In such cells the copy number of a ColE1-derived plasmid is decreased. The cell may further comprise an expression element comprising an inducible promoter operably linked to an ORF encoding the wild-type *pcnB*. Because the wild-type *pcnB* gene is dominant to the mutant *pcnB80* gene, and because the wild-type *pcnB* gene product promotes increased ColE1 plasmid copy number, induction of a wild-type *pcnB* in the *pcnB80* background will increase the plasmid copy number of ColE1-derived plasmids. Such copy number control systems may be expressed from the chromosome and/or plasmid to maintain either low or high plasmid copy number in the absence of induction. Other non-limiting examples of gene and/or gene products that may be employed in copy number control systems for ColE1-based replicons include genes or homologs of genes encoding RNAI, RNA II, *rop*, RNase H, enzymes involved in the process of polyadenylation, RNase E, DNA polymerase I, and DNA polymerase III.

[0361] In the case of IncFII-derived replicons, non-limiting examples of gene and/or gene products that may be employed in copy number control systems to control plasmid copy include genes or homologs of the *copA*, *copB*, *repA*, and *repB* genes. Copy number control systems may additionally or alternatively include manipulation of *repC*, *trfA*, *dnaA*, *dnaB*, *dnaC*, *seqA*, genes protein Pi, genes encoding HU protein subunits (*hupA*, *hupB*) and genes encoding IHF subunits.

[0362] Other elements may also be included to optimize these plasmid copy number control systems. Such additional elements may include the addition or deletion of iteron nucleic acid sequences (Chattoraj, D. K. 2000. Control of plasmid DNA replication by iterons: no longer paradoxical. *Mol. Microbiol.* 37:467-476), and modification of chaperone proteins involved in plasmid replication (Konieczny, I., et al. 1997. The replication initiation protein of the broad-host-range plasmid RK2 is activated by the ClpX chaperone. *Proc Natl Acad Sci USA* 94:14378-14382).

#### [0363] II.C.6. Transportation of Inducible and Inhibitory Compounds

[0364] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include utilization and/or modification of factors and systems that modulate the transport of compounds, including but not limited to inducers and/or inhibitors of expression elements that control expres-

sion of a gene in a parent cell prior to minicell formation and/or in segregated minicells. Such manipulations may result in increased or decreased production, and/or changes in the intramolecular and intermolecular functions, of a protein in a minicell or its parent cell. The techniques may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells.

**[0365]** II.C.6.a. *Escherichia coli* Genes

**[0366]** By way of non-limiting example, manipulation of the *abpS* gene or gene product from *E. coli*, or homologs of this gene or gene product found in other members of the Prokaryotes, Eukaryotes, Archaeobacteria and/or organelles (e.g., mitochondria, chloroplasts, plastids and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells (Celis, R. T. 1982. Mapping of two loci affecting the synthesis and structure of a periplasmic protein involved in arginine and ornithine transport in *Escherichia coli* K-12. *J. Bacteriol.* 151(3):1314-9).

**[0367]** In addition to *abpS*, other exemplary *E. coli* genes encoding factors involved in the transport of inducers, inhibitors and other compounds include, but are not limited to, *araE* (Khlebnikov, A., et al. 2001. Homogeneous expression of the P(BAD) promoter in *Escherichia coli* by constitutive expression of the low-affinity high-capacity AraE transporter. *Microbiology.* 147(Pt 12):3241-7); *araG* (Kehres, D. G., and Hogg, R. W. 1992. *Escherichia coli* K12 arabinose-binding protein mutants with altered transport properties. *Protein Sci.* 1(12): 1652-60); *araH* (Id.); *argP* (Celis, R. T. 1999. Repression and activation of arginine transport genes in *Escherichia coli* K 12 by the ArgP protein. *J. Mol. Biol.* 17;294(5):1087-95); *aroT* (*aroR*, *trpR*) (Edwards, R. M., and Yudkin, M. D. 1982. Location of the gene for the low-affinity tryptophan-specific permease of *Escherichia coli*. *Biochem. J.* 204(2):617-9); *artI* (Wissenbach, U., et al. 1995. A third periplasmic transport system for L-arginine in *Escherichia coli*: molecular characterization of the *artIQMJ* genes, arginine binding and transport. *Mol. Microbiol.* 17(4):675-86); *artJ* (Id.); *artM* (Id.); *artP* (Id.); *artQ* (Id.); *bioP* (*bir*, *birB*) (Campbell, A., et al. Biotin regulatory (*bir*) mutations of *Escherichia coli*. 1980. *J. Bacteriol.* 142(3):1025-8); *brnQ* (*hrbA*) (Yamato, I., and Anraku, Y. 1980. Genetic and biochemical studies of transport systems for branched-chain amino acids in *Escherichia coli* K-12: isolation and properties of mutants defective in leucine-repressible transport activities. *J. Bacteriol.* 144(1):36-44); *brnR* (Id.); *brnS* (Id.); *brnT* (Id.); *btuC* (Friedrich, M. J., et al. 1986. Nucleotide sequence of the *btuCED* genes involved in vitamin B12 transport in *Escherichia coli* and homology with components of periplasmic-binding-protein-dependent transport systems. *J. Bacteriol.* 167(3):928-34); *btuD* (Id.) (Friedrich, M. J., et al. 1986. Nucleotide sequence of the *btuCED* genes involved in vitamin B12 transport in *Escherichia coli* and homology with components of periplasmic-binding-protein-dependent transport systems. *J. Bacteriol.* 167(3):928-34); *caiT* (Eichler, K. 1994. Molecular characterization of the *cai* operon necessary for carnitine metabolism in *Escherichia coli*. *Mol. Microbiol.* 13(5):775-86); *celA* (Parker, L. L., and Hall, B. G. 1990. Characterization and nucleotide sequence of the cryptic *cel* operon of *Escherichia coli* K12. *Genetics.*

124(3):455-71); *celB* (Id.); *celC* (Id.); *citA* (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein); *citB* (Id.); *codB* (Danielsen, S., et al. 1992. Characterization of the *Escherichia coli* *codBA* operon encoding cytosine permease and cytosine deaminase. *Mol. Microbiol.* 6(10):1335-44); *cysA* (Karbonowska, H., et al. 1977. Sulphate permease of *Escherichia coli* K12. *Acta. Biochim. Pol.* 24(4):329-34); *cysU* (*cyst*) (Sirko, A., et al. 1995. Sulfate and thiosulfate transport in *Escherichia coli* K-12: evidence for a functional overlapping of sulfate- and thiosulfate-binding proteins. *J. Bacteriol.* 177(14):4134-6); *cysW* (Id.); *dctA* (Lo, T. C., and Bewick, M. A. 1978. The molecular mechanisms of dicarboxylic acid transport in *Escherichia coli* K12. The role and orientation of the two membrane-bound dicarboxylate binding proteins. *J. Biol. Chem.* 10;253(21):7826-31); *dctB* (Id.); *dcuA* (*genA*) (Six, S., et al. 1994. *Escherichia coli* possesses two homologous anaerobic C4-dicarboxylate membrane transporters (*DcuA* and *DcuB*) distinct from the aerobic dicarboxylate transport system (*Dct*). *J. Bacteriol.* 176(21):6470-8); *dcuB* (*genF*) ( ); *dgoT* (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein); *exuT* (Nemoz, G., et al. 1976. Physiological and genetic regulation of the aldohexuronate transport system in *Escherichia coli*. *J. Bacteriol.* 127(2):706-18); *fepD* (Ozenberger, B. A., et al. 1987. Genetic organization of multiple *fep* genes encoding ferric enterobactin transport functions in *Escherichia coli*. *J. Bacteriol.* 169(8):3638-46); *fepG* (Chenault, S. S., and Earhart, C. F. 1991. Organization of genes encoding membrane proteins of the *Escherichia coli* ferrienterobactin permease. *Mol. Microbiol.* 5(6):1405-13); *fucP* (*prd*) (Chen, Y. M. 1987. The organization of the *fuc* regulon specifying L-fucose dissimilation in *Escherichia coli* K12 as determined by gene cloning. *Mol. Gen. Genet.* 210(2):331-7); *glnP* (Masters, P. S., and Hong, J. S. 1981. Genetics of the glutamine transport system in *Escherichia coli*. *J. Bacteriol.* 147(3):805-19); *glnQ* (Nohno, T. 1986. Cloning and complete nucleotide sequence of the *Escherichia coli* glutamine permease operon (*glnHPQ*). *Mol. Gen. Genet.* 205(2):260-9); *glnR* (Masters, P. S., and Hong, J. S. 1981. Genetics of the glutamine transport system in *Escherichia coli*. *J. Bacteriol.* 147(3):805-19); *glpT* (Boos, W., et al. 1977. Purification and properties of a periplasmic protein related to sn-glycerol-3-phosphate transport in *Escherichia coli*. *Eur. J. Biochem.* 72(3):571-81); *glpP* (Deguchi, Y., et al. 1989. Molecular cloning of *gltS* and *gltP*, which encode glutamate carriers of *Escherichia coli*. *B. J. Bacteriol.* 171(3):1314-9); *gltS* (Id.); *gntR* (Bachi, B., and Kornberg, H. L. 1975. Genes involved in the uptake and catabolism of gluconate by *Escherichia coli*. *J. Gen. Microbiol.* 90(2):321-35); *gntS* (Id.); *gntT* (*gntM*, *usgA*) (Id.); *gntU* (Tong, S. 1996. Cloning and molecular genetic characterization of the *Escherichia coli* *gntR*, *gntK*, and *gntU* genes of *GntI*, the main system for gluconate metabolism. *J. Bacteriol.* 178(11):3260-9); *hisM* (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella*



- typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein); his P (Id.); his Q (Id.); livG (hrbB, hrbC, hrbD) (Landick, R., et al. 1980. Regulation of high-affinity leucine transport in *Escherichia coli*. J. Supramol. Struct. 14(4):527-37); livH (hrbB, hrbC, hrbD) (Id.); livJ (hrbB, hrbC, hrbD) (Id.); livK (hrbB, hrbC, hrbD) (Id.); livM (Id.); lldP (letP) (Dong, J. M., et al. 1993. Three overlapping let genes involved in L-lactate utilization by *Escherichia coli*. J. Bacteriol. 175(20):6671-8); lysP (cadR) (Steffes, C., et al. 1992. The lysP gene encodes the lysine-specific permease. J. Bacteriol. 174(10):3242-9); malF (malB) (Bavoil, P., et al. 1980. Identification of a cytoplasmic membrane-associated component of the maltose transport system of *Escherichia coli*. J. Biol. Chem. 255(18):8366-9); malG (malB) (Dassa, E., and Hofnung, M. 1985. Sequence of gene malG in *E. coli* K12: homologies between integral membrane components from binding protein-dependent transport systems. EMBO J. 4(9):2287-93); malK (malB) (Id.); mglC (PMG, mglP) (Harayama, S. 0.1983. Characterization of the mgl operon of *Escherichia coli* by transposon mutagenesis and molecular cloning. J. Bacteriol. 153(1):408-15); nanT (Vimr, E. R., and Troy, F. A. 1985. Identification of an inducible catabolic system for sialic acids (nan) in *Escherichia coli*. J. Bacteriol. 164(2):845-53); nupC (cru) (Craig, J. E., et al. 1994. Cloning of the nupC gene of *Escherichia coli* encoding a nucleoside transport system, and identification of an adjacent insertion element, IS 186. Mol. Microbiol. 11(6):1159-68); nupG (Westh Hansen, S. E., et al. 1987. Studies on the sequence and structure of the *Escherichia coli* K-12 nupG gene, encoding a nucleoside-transport system. Eur. J. Biochem. 168(2):385-91); panF (Vallari, D. S., and Rock, C. O. 1985. Isolation and characterization of *Escherichia coli* pantothenate permease (panF) mutants. J. Bacteriol. 164(1):136-42); potA (Kashiwagi, K., et al. 1993. Functions of potA and potD proteins in spermidine-preferential uptake system in *Escherichia coli*. J. Biol. Chem. 268(26): 19358-63); potG (Pistocchi, R., et al. 1993. Characteristics of the operon for a putrescine transport system that maps at 19 minutes on the *Escherichia coli* chromosome. J. Biol. Chem. 268(1):146-52); potH (Id.); potI (Id.); proP (Wood, J. M., and Zadworny, D. 1980. Amplification of the put genes and identification of the put gene products in *Escherichia coli* K12. Can. J. Biochem. 58(10):787-96); proT (Id.); proV (proU) (Faatz, E., et al. 1988. Cloned structural genes for the osmotically regulated binding-protein-dependent glycine betaine transport system (ProU) of *Escherichia coli* K-12. Mol. Microbiol. 2(2):265-79); proW (proU) (Id.); proX (proU) (Id.); pstA (R2pho, phoR2b, phoT) (Amemura, M., et al. 1985. Nucleotide sequence of the genes involved in phosphate transport and regulation of the phosphate regulon in *Escherichia coli*. J. Mol. Biol. 184(2):241-50); pstB (phoT) (Id.); pstC (phoW) (Rao, N. N., and Torriani, A. 1990. Molecular aspects of phosphate transport in *Escherichia coli*. Mol. Microbiol. 4(7):1083-90); pstS (R2pho, nmpA, phoR2a, phoS) (Makino, K., et al. 1988. Regulation of the phosphate regulon of *Escherichia coli*. Activation of pstS transcription by PhoB protein in vitro. J. Mol. Biol. 203(1):85-95); purP (Burton, K. 1994. Adenine transport in *Escherichia coli*. Proc. R. Soc. Lond. B. Biol. Sci. 255(1343):153-7); putP (Stalmach, M. E., et al. 1983. Two proline porters in *Escherichia coli* K-12. J. Bacteriol. 156(2):481-6); rbsA (rbsP, rbsT) (Iida, A., et al. 1984. Molecular cloning and characterization of genes required for ribose transport and utilization in *Escherichia coli* K-12. J. Bacteriol. 158(2):674-82); rbsC (rbsP, rbsT) (Id.); rbsD (rbsP) (Id.); rhaT (Baldoma, L., et al. 1990. Cloning, mapping and gene product identification of rhaT from *Escherichia coli* K12. FEMS Microbiol. Lett. 60(1-2):103-7); sdaC (Shao, Z., et al. 1994. Sequencing and characterization of the sdaC gene and identification of the sdaCB operon in *Escherichia coli* K12. Eur. J. Biochem. 222(3):901-7); tnaB (trpP) (Sarsero, J. P., et al. 1991. A new family of integral membrane proteins involved in transport of aromatic amino acids in *Escherichia coli*. J. Bacteriol. 173(10):3231-4); tyrR (Whipp, M. J., and Pittard, A. J. 1977. Regulation of aromatic amino acid transport systems in *Escherichia coli* K-12. J. Bacteriol. 132(2):453-61); ugpC (Schweizer, H., and Boos, W. 1984. Characterization of the ugp region containing the genes for the phoB dependent sn-glycerol-3-phosphate transport system of *Escherichia coli*. Mol. Gen. Genet. 197(1): 161-8); uhpT (Weston, L. A., and Kadner, R. J. 1987. Identification of uhp polypeptides and evidence for their role in exogenous induction of the sugar phosphate transport system of *Escherichia coli* K-12. J. Bacteriol. 169(8):3546-55); and xylF (xylT) (Sumiya, M., et al. 1995. Molecular genetics of a receptor protein for D-xylose, encoded by the gene xylF, in *Escherichia coli*. Receptors Channels. 3(2):117-28).
- [0368]** II.C.6.b. *Bacillus subtilis* Genes
- [0369]** By way of non-limiting example, manipulation of the aapA gene or gene product from *B. subtilis*, or homologs of this gene or gene product found in other members of the Prokaryotes, Eukaryotes, Archaeobacteria and/or organelles (e.g., mitochondria, chloroplasts, plastids and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.).
- [0370]** In addition to aapA, other exemplary *B. subtilis* genes encoding factors involved in the transport of inducers, inhibitors and other compounds include, but are not limited to, arnyC (Sekiguchi, J., et al. 1975. Genes affecting the productivity of alpha-amylase in *Bacillus subtilis*. J. Bacteriol. 121(2):688-94); amyD (Id.); araE (Sa-Nogueira, I., and Mota, L. J. 1997. Negative regulation of L-arabinose metabolism in *Bacillus subtilis*: characterization of the araR (araC) gene. J. Bacteriol. 179(5): 1598-608); araN (Sa-Nogueira, I., et al. 1997. The *Bacillus subtilis* L-arabinose (ara) operon: nucleotide sequence, genetic organization and expression. Microbiology. 143 (Pt 3):957-69); araP (Id.); araQ (Id.); csbC (Akbar, S., et al. 1999. Two genes from *Bacillus subtilis* under the sole control of the general stress transcription factor sigmaB. Microbiology. 145 (Pt 5):1069-78); cysP (Mansilla, M. C., and de Mendoza, D. 2000. The *Bacillus subtilis* cysP gene encodes a novel sulphate permease related to the inorganic phosphate transporter (Pit) family. Microbiology. 146 (Pt 4):815-21); dctB (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); exuT (Rivolta, C., et al. 1998. A 35.7 kb DNA fragment from the *Bacillus subtilis* chromosome containing a putative 12.3 kb operon



involved in hexuronate catabolism and a perfectly symmetrical hypothetical catabolite-responsive element. *Microbiology*. 144 (Pt 4):877-84); gabP (Ferson, A. E., et al. 1996. Expression of the *Bacillus subtilis* gabP gene is regulated independently in response to nitrogen and amino acid availability. *Mol. Microbiol.* 22(4):693-701); gamp (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); glcP (Paulsen, I. T., et al. 1998. Characterization of glucose-specific catabolite repression-resistant mutants of *Bacillus subtilis*: identification of a novel hexose:H<sup>+</sup> symporter. *J. Bacteriol.* 180(3):498-504); glcU (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); glnH (Id.); glnM (Id.); glnP (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); glnQ (Id.); glpT (Nilsson, R. P., et al. 1994. The glpT and glpQ genes of the glycerol regulon in *Bacillus subtilis*. *Microbiology*. 140 (Pt 4):723-30); gltP (Tolner, B., et al. 1995. Characterization of the proton/glutamate symport protein of *Bacillus subtilis* and its functional expression in *Escherichia coli*. *J. Bacteriol.* 177(10):2863-9); gltT (Tolner, B., et al. 1995. Characterization of the proton/glutamate symport protein of *Bacillus subtilis* and its functional expression in *Escherichia coli*. *J. Bacteriol.* 177(10):2863-9); gntP (Reizer, A., et al. Analysis of the gluconate (gnt) operon of *Bacillus subtilis*. *Mol. Microbiol.* 5(5):1081-9); gutP (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); hutM (Oda, M., et al. 1988. Cloning and nucleotide sequences of histidase and regulatory genes in the *Bacillus subtilis* hut operon and positive regulation of the operon. *J. Bacteriol.* 170(7):3199-205); iofP (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); kdgT (Pujic, P., et al. 1998. The kdgRKAT operon of *Bacillus subtilis*: detection of the transcript and regulation by the kdgR and ccpA genes. *Microbiology*. 144 (Pt 11):3111-8); lctP (Cruz, Ramos H., et al. 2000. Fermentative metabolism of *Bacillus subtilis*: physiology and regulation of gene expression. *J. Bacteriol.* 182(11):3072-80); maeN (Ito, M., et al. 2000. Effects of nonpolar mutations in each of the seven *Bacillus subtilis* mrp genes suggest complex interactions among the gene products in support of Na<sup>+</sup> and alkali but not cholera resistance. *J. Bacteriol.* 182(20):5663-70); malP (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); manP (Id.); mleN (Id.); nasA (Ogawa, K., et al. 1995. The nasB operon and nasA gene are required for nitrate/nitrite assimilation in *Bacillus subtilis*. *J. Bacteriol.* 177(5):1409-13); nupC (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); opuAB (Kempf, B., et al. 1997. Lipoprotein from the osmoregulated ABC transport system OpuA of *Bacillus subtilis*: purification of the glycine betaine binding protein and characterization of a functional lipidless mutant. *J. Bacteriol.* 179(20):6213-20); opuBA (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); pbuG (Saxild, H. H., et al. 2001. Definition of the *Bacillus subtilis* PurR operator using genetic and bioinformatic tools and expansion of the PurR regulon with glyA, guaC, pbuG, xpt-pbuX, yqhZ-fofD, and pbuO. *J. Bacteriol.* 183(21):6175-83); pbuX (Saxild, H. H., et al. 2001. Definition of the *Bacillus subtilis* PurR operator using genetic and bioinformatic tools and expansion of the PurR regulon with glyA, guaC, pbuG, xpt-pbuX, yqhZ-fofD, and pbuO. *J. Bacteriol.* 183(21):6175-83); pstC (Takemaru, K., et al. 1996. A *Bacillus subtilis* gene cluster similar to the *Escherichia coli* phosphate-specific transport (pst) operon: evidence for a tandemly arranged pstB gene. *Microbiology*. 142 (Pt 8):2017-20); pstS (Qi, Y., et al. 1997. The pst operon of *Bacillus subtilis* has a phosphate-regulated promoter and is involved in phosphate transport but not in regulation of the pho regulon. *J. Bacteriol.* 179(8):2534-9); pucJ (Schultz, A. C., et al. 2001. Functional analysis of 14 genes that constitute the purine catabolic pathway in *Bacillus subtilis* and evidence for a novel regulon controlled by the PucR transcription activator. *J. Bacteriol.* 183(11):3293-302); pucK (Schultz, A. C., et al. 2001. Functional analysis of 14 genes that constitute the purine catabolic pathway in *Bacillus subtilis* and evidence for a novel regulon controlled by the PucR transcription activator. *J. Bacteriol.* 183(11):3293-302); pyrP (Turner, R. J., et al. 1994. Regulation of the *Bacillus subtilis* pyrimidine biosynthetic (pyr) gene cluster by an autogenous transcriptional attenuation mechanism. *J. Bacteriol.* 176(12):3708-22); rbsB (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); rbsC (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); rbsD (Id.); rocC (Gardan, R., et al. 1995. Expression of the rocDEF operon involved in arginine catabolism in *Bacillus subtilis*. *J. Mol. Biol.* 23;249(5):843-56); rocE (Gardan, R., et al. 1995. Expression of the rocDEF operon involved in arginine catabolism in *Bacillus subtilis*. *J. Mol. Biol.* 23;249(5):843-56); ssuA (Coppee, J. Y., et al. 2001. Sulfur-limitation-regulated proteins in *Bacillus subtilis*: a two-dimensional gel electrophoresis study. *Microbiology*. 147(Pt 6):1631-40); ssuB (van der Ploeg, J. R., et al. 1998. *Bacillus subtilis* genes for the utilization of sulfur from aliphatic sulfonates. *Microbiology*. 144 (Pt 9):2555-61); ssuC (van der Ploeg, J. R., et al. 1998. *Bacillus subtilis* genes for the utilization of sulfur from aliphatic sulfonates. *Microbiology*. 144 (Pt 9):2555-61); treP (Yamamoto, H., et al. 1996. Cloning and sequencing of a 40.6 kb segment in the 73 degrees-76 degrees region of the *Bacillus subtilis* chromosome containing genes for trehalose metabolism and acetoin utilization. *Microbiology*. 142 (Pt 11):3057-65); xynP (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); ybaR (Id.); ybgF (Id.); ybgH (Id.); ycbE (Id.); ycgO (Id.); yckI (Id.); yckJ (Id.); yckK (Id.); ydgF (Id.); yecA (Borriess, R., et al. 1996. The 52 degrees-55 degrees segment of the *Bacillus subtilis* chromosome: a region devoted to purine uptake and metabolism, and containing the genes cotA, gabP and guaA and the pur gene cluster within a 34960 bp nucleotide sequence. *Microbiology*. 142 (Pt 11):3027-31); yesP (Sohenshein, A. L., J. A.

Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); yesQ (Id.); yfIS (Id.); yhcL (Id.); yhjB (Id.); yjkB (Id.); ykbA (Id.); yoaB (Id.); yocN (Id.); yodF (Id.); yojA (Id.); yqiY (Id.); ytlD (Id.); ytlP (Id.); ytmL (Id.); ytmM (Id.); ytnA (Id.); yurM (Id.); yurN (Id.); yvbW (Id.); yvdH (Id.); yvdI (Id.); yveA (Pereira, Y., et al. 2001. The yveB gene, Encoding endolevanase LevB, is part of the sacB-yveB-yveA levansucrase tricistronic operon in *Bacillus subtilis*. Microbiology. 147(Pt 12):3413-9); yvfH (Sohenshein, A. L., J. A. Hoch, and R. Losick (eds.) 2002. *Bacillus subtilis* and its closest relatives: from genes to cells. American Society for Microbiology, Washington D.C.); yvFL (Id.); yvFM (Id.); yvgM (Id.); yvrO (Id.); yvsH (Id.); ywbF (Id.); ywcJ (Id.); ywoD (Id.); ywoE (Id.); yxeN (Id.); and yxeR (Id.).

#### [0371] II.C.7. Catabolite Repression

[0372] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include utilization and/or modification of factors and systems involved in the synthesis, degradation or transport of catabolites that modulate the genetic expression of a preselected protein. Such manipulations may result in increased or decreased production, and/or changes in the intramolecular and intermolecular functions, of a protein in a minicell or its parent cell; in the latter instance, the protein may be one that is desirably retained in segregated minicells.

[0373] By way of non-limiting example, it is known in the art to use promoters from the trp, cst-1, and llp operons of *E. coli*, which are induced by, respectively, reduced tryptophan levels, glucose starvation, and lactose. Manipulation of the catabolites tryptophan, glucose and lactose, respectively, will influence the degree of expression of genes operably linked to these promoters. (Makrides, Savvas C., Strategies for Achieving High-Level Expression of Genes in *Escherichia coli*. Microbiological Reviews. 1996. 60:512-538.)

[0374] As another non-limiting example, expression elements from the *E. coli* L-arabinose (ara) operon are used in expression systems. AraC is a protein that acts as a repressor of ara genes in the absence of arabinose, and also as an activator of ara genes when arabinose is present. Induction of ara genes also involves cAMP, which modulates the activity of CRP (cAMP receptor protein), which in turn is required for full induction of ara genes (Schleif, Robert, Regulation of the L-arabinose operon of *Escherichia coli*. 2000. TIG 16:559-564. Thus, maximum expression from an ara-based expression system is achieved by adding cAMP and arabinose to host cells, and optimizing the expression of CRP in hostcells.

[0375] As one example, manipulation of the acpS gene or gene product of *E. coli* (Pollacco M. L., and J. E. Cronan Jr. 1981. A mutant of *Escherichia coli* conditionally defective in the synthesis of holo-[acyl carrier protein]. J. Biol.Chem. 256:5750-5754); or homologs of this gene or its gene product found in other prokaryotes, eukaryotes, archaeobacteria or organelles (mitochondria, chloroplasts, plastids and the like) may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or in segregated minicells.

[0376] In addition to acpS, other exemplary *E. coli* genes include the b2383 gene (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein. b2387 gene; the celA gene (Parker L. L., and B. G. Hall. 1990. Characterization and nucleotide sequence of the cryptic cel operon of *Escherichia coli* K12. Genetics. 124:455-471); the celB gene (Cole S. T., and B. Saint-Joanis, and A. P. Pugsley. 1985. Molecular characterisation of the colicin E2 operon and identification of its products. Mol Gen Genet. 198:465-472); the celC gene (Parker L. L., and B. G. Hall. 1990. Characterization and nucleotide sequence of the cryptic cel operon of *Escherichia coli* K12. Genetics. 124:455-471); the cmtB gene (Ezhova N. M., Zaikina, N. A, Shataeva, L. K., Dubinina, N. I., Ovechkina, T. P. and J. V. Kopylova. [Sorption properties of carboxyl cation exchangers with a bacteriostatic effect]. 1980. Prikl Bioikhim Mikrobiol. 16:395-398); the creB gene (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein; the creC gene (Wanner B. L. Gene regulation by phosphate in enteric bacteria. 1993. J Cell Biochem. 51:47-54); the crp gene (Sabourn D., and J. Beckwith. Deletion of the *Escherichia coli* crp gene. 1975. J Bacteriology. 122:338-340); the crr (gsr, iex, tgs, treD) gene (Jones-Mortimer M. C., and H. L. Kornberg, and r. Maltby, and P. D. Watts. Role of the crr-gene in glucose uptake by *Escherichia coli*. 1977. FEBS Lett. 74:17-19); the cya gene (Bachi B., and H. L. Kornberg. Utilization of gluconate by *Escherichia coli*. A role of adenosine 3':5'-cyclic monophosphate in the induction of gluconate catabolism. 1975. Biochem J. 150:123-128); thefruA gene (Prior T. I., and H. L. Kornberg. Nucleotide sequence of fruA, the gene specifying enzyme Iifru of the phosphoenopyruvate-dependent sugar phosphotransferase system in *Escherichia coli* K12. 1988. J Gen Microbiol. 134:2757-2768); the fruB gene (Bol'shakova T. N. and R. S. Erlagaeva, and Dobrynya Oiu, and V. N. Gershanovich. [Mutation fruB in the fructose regulon affecting beta-galactosidase synthesis and adenylate cyclase activity of *E. coli* K12]. 1988. Mol Gen Mikrobiol virusol. 3:33-39); thefruR gene (Jahreis K., and P. W. Postma, and J. W. Lengeler. Nucleotide sequence of the ilvH-frR gene region of *Escherichia coli* K12 and *Salmonella typhimurium* LT2. 1991. Mol Gen Genet. 226:332-336); the frvA gene (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein); the ftwB gene (Id.); the frvD gene (Id.); the gatB gene (Nobelmann B., and J. W. Lengeler. Molecular analysis of the gat genes from *Escherichia coli* and of their roles in galactitol transport and metabolism. 1996. J Bacteriol. 178:6790-6795); the gatC gene (Id.); the malX gene (Reidel J., W. Boos. The malX malY operon of *Escherichia coli* encodes a novel enzyme II of the photophototransferase system recognizing glucose and maltose and an enzyme

abolishing the endogenous induction of the maltose system. 1991. *J. Bacteriol.* 173:4862-4876); the manX (gptB, mpt, ptsL, ptsM, ptsX, manIII) gene (Plumbridge J., and A. Kolb. CAP and Nag repressor binding to the regulatory regions of the nagE-B and manX genes of *Escherichia coli*. 1991. *J. Mol. Biol.* 217:661-679); the manY (pel, ptsM, ptsP, manPII) gene (Henderson P. J., and R. A. Giddens, and M. C. Jones-Mortimer. Transport of galactose, glucose and their molecular analogues by *Escherichia coli* K12. 1977. *Biochem J.* 162:309-320); the manZ (gptB, mpt, ptsM, ptsX) gene (Williams N., and D. K. Fox, and C. Shea and S. Roseman. Pel, the protein that permits lambda DNA penetration of *Escherichia coli*, is encoded by a gene in ptsM and is required for mannose utilization by the phosphotransferase system. 1986. *Proc Natl Acad Sci USA.* 83:8934-8938); the mtlA gene (Lengeler J. Mutations affecting transport of the hexitols D-mannitol, D-glucitol, and galactitol in *Escherichia coli* K-12: isolation and mapping. 1975. *J. Bacteriol.* 124:26-38.); the nagE (pstn) gene (Rogers M. J., and T. Ohgi, and J. Plumbridge, and D. Soll. Nucleotide sequences of the *Escherichia coli* nagE and nagB genes: the structural genes for the N-acetylglucosamine transport protein of the bacterial phosphoenolpyruvate: sugar phosphotransferase system and for glucosamine-6-phosphate deaminase. 1988. *Gene.* 62:197-207); the pstA gene (Cox G. B., H. Rosenberg, and J. A. Downie, and S. Silver. Genetic analysis of mutants affected in the Pst inorganic phosphate transport system. 1981. *J. Bacteriol.* 148:1-9); the pstB (gutB) gene (Id.); the pstG gene (Cox G. B., H. Rosenberg, and J. A. Downie, and S. Silver. Genetic analysis of mutants affected in the Pst inorganic phosphate transport system. 1981. *J. Bacteriol.* 148:1-9); the pstH gene (Id.); the pstI gene (Id.); the pstN gene (Id.); the pstO gene (Id.); the ptxA (yifU) gene (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein); the sgcA (yjhL) gene (Id.); the sgcC (yjhN) gene (Id.); the treB gene (Boos W., U. Ehmann, H. Forkl, W. Klein, M. Rimmele, and P. Postma. Trehalose transport and metabolism in *Escherichia coli*. 1990. *J. Bacteriol.* 172:3450-3461); the usg gene (Arps P. J., and M. E. Winkler M E. Structural analysis of the *Escherichia coli* K-12 hisT operon by using a kanamycin resistance cassette. 1987. *J. Bacteriol.* 169:1061-1070); the wcaD gene (Mao Y., and M. P. Doyle, and J. Chen. Insertion mutagenesis of wca reduces acid and heat tolerance of enterohemorrhagic *Escherichia coli* O157:H7. 2001. *J. Bacteriol.* 183:3811-3815); the yadL gene (Berlyn et al., "Linkage Map of *Escherichia coli* K-12, Edition 9," Chapter 109 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1715-1902, and references cited therein); and the ycgC gene (Gutknecht R., and R. Beutler, and L. F. Garcia-Alles, and U. Baumann, and B. Erni. The dihydroxyacetone kinase of *Escherichia coli* utilizes a phosphoprotein instead of ATP as phosphoryl donor. 2001. *EMBO J.* 20:2480-2486).

#### [0377] II.C.8. General Deletions and Modifications

[0378] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting

example, these techniques may include modification or deletion of endogenous gene(s) from which their respective gene product decreases the induction and expression efficiency of a desired protein in the parent cell prior to minicell formation and/or the segregated minicell. By way of non-limiting example, these protein components may be the enzymes that degrade chemical inducers of expression, proteins that have a dominant negative affect upon a positive regulatory elements, proteins that have proteolytic activity against the protein to be expressed, proteins that have a negative affect against a chaperone that is required for proper activity of the expressed protein, and/or this protein may have a positive effect upon a protein that either degrades or prevents the proper function of the expressed protein. These gene products that require deletion or modification for optimal protein expression and/or function may also be antisense nucleic acids that have a negative affect upon gene expression.

#### [0379] II.C.9. Cytoplasmic Redox State

[0380] Included in the design of the invention are techniques that increase the efficiency of gene expression and functional protein production in minicells. By way of non-limiting example, these techniques may include modification of endogenous and/or exogenous protein components that alter the redox state of the parental cell cytoplasm prior to minicell formation and/or the segregated minicell cytoplasm. By way of non-limiting example, this protein component may be the product of the *trxA*, *grx*, *dsbA*, *dsbB*, and/or *dsbc* genes from *E. coli* or homologs of this gene or gene product found in other members of the Eubacteria, Eucarya or Archae (Mark et al., Genetic mapping of *trxA*, a gene affecting thioredoxin in *Escherichia coli* K12, *Mol Gen Genet.* 155:145-152, 1977; (Russel et al., Thioredoxin or glutaredoxin in *Escherichia coli* is essential for sulfate reduction but not for deoxyribonucleotide synthesis, *J. Bacteriol.* 172:1923-1929, 1990); Akiyama et al., In vitro catalysis of oxidative folding of disulfide-bonded proteins by the *Escherichia coli* *dsbA* (*ppfA*) gene product, *J. Biol. Chem.* 267:22440-22445, 1992); (Whitney et al., The DsbA-DsbB system affects the formation of disulfide bonds in periplasmic but not in intramembraneous protein domains, *FEBS Lett.* 332:49-51, 1993); (Shevchik et al., Characterization of DsbC, a periplasmic protein of *Erwinia chrysanthemi* and *Escherichia coli* with disulfide isomerase activity, *EMB J.* 13:2007-2012, 1994). These applications may, but are not limited to increased or decreased production, increased or decreased intramolecular TrxA activity, increased or decreased physiological function of the above-mentioned gene products. By way of non-limiting example, increased production of gene product (gene expression) may occur through increased gene dosage (increased copy number of a given gene under the control of the native or artificial promoter where this gene may be on a plasmid or in more than one copy on the chromosome), modification of the native regulatory elements, including, but not limited to the promoter or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/inhancers, or relevant antisense nucleic acid or nucleic acid analog, cloning on a plasmid under the control of the native or artificial promoter, and increased or decreased production of native or artificial promoter regulatory elements) controlling production of the gene. By way of non-limiting example, decreased gene expression production may occur through modification of the native regulatory

elements, including, but not limited to the promoter or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/inhancers, or relevant antisense nucleic acid or nucleic acid analog, through cloning on a plasmid under the control of the native regulatory region containing mutations or an artificial promoter, either or both of which resulting in decrease gene expression, and through increased or decreased production of native or artificial promoter regulatory element(s) controlling gene expression. By definition, intramolecular activity refers to the enzymatic function, structure-dependent function, e.g. the capacity of a gene product to interact in a protein-protein, protein-nucleic acid, or protein-lipid complex, and/or carrier function, e.g. the capacity to bind, either covalently or non-covalently small organic or inorganic molecules, protein(s) carbohydrate(s), fatty acid(s), lipid(s), and nucleic acid(s). By way of non-limiting example, alteration of intramolecular activity may be accomplished by mutation of the gene, in vivo or in vitro chemical modification of the gene product, inhibitor molecules against the function of the gene product, e.g. competitive, non-competitive, or uncompetitive enzymatic inhibitors, inhibitors that prevent protein-protein, protein-nucleic acid, or protein-lipid interactions, e.g. expression or introduction of dominant-negative or dominant-positive or other protein fragment(s), or other carbohydrate(s), fatty acid(s), lipid(s), and nucleic acid(s) that may act directly or allosterically upon the gene product, and/or modification of protein, carbohydrate, fatty acid, lipid, or nucleic acid moieties that modify the gene or gene product to create the functional protein. By definition, physiological function refers to the effects resulting from an intramolecular interaction between the gene product and other protein, carbohydrate, fatty acid, lipid, nucleic acid, or other molecule(s) in or on the cell or the action of a product or products resulting from such an interaction.

[0381] By way of non-limiting example, physiological function may be the act or result of intermolecular phosphorylation, biotinylation, methylation, acylation, glycosylation, and/or other signaling event; this function may be the result of protein-protein, protein-nucleic acid, or protein-lipid interaction resulting in a functional moiety; this function may be to interact with the membrane to recruit other molecules to this compartment of the cell; this function may be to regulate the transcription and/or translation of trxA, other protein, or nucleic acid; and this function may be to stimulate the function of another process that is not yet described or understood.

[0382] II.C.10. Transcriptional Terminators

[0383] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in parental cell cytoplasm prior to minicell formation and/or the segregated minicell cytoplasm. By way of non-limiting example, these techniques may include modification of terminator regions of DNA templates or RNA transcripts so that transcription and/or translation of these nucleic acid regions will terminate at greater efficiency. By way of non-limiting example, these techniques may include stem-loop structures, consecutive translational terminators, polyadenylation sequences, or increasing the efficiency of rho-dependent termination. Stem loop structures may include, but are not limited to, inverted repeats containing any combination of deoxyribonucleic acid or

ribonucleic acid molecule, more than one such inverted repeat, or variable inverted repeats such that the rate of transcriptional/translational termination may be moderated dependent on nucleic acid and/or amino acid concentration, e.g. the mechanism of regulatory attenuation (Oxender et al., Attenuation in the *Escherichia coli* tryptophan operon: role of RNA secondary structure involving the tryptophan codon region, Proc. Natl. Acad. Sci. 76:5524-5528, 1979). See also, Yager and von Hippel, "Transcript Elongation and Termination in e. Col. And Landick and Yanofsky, "Transcriptional Attenuation," Chapters 76 and 77, respectively in: *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*, Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1987, Volume 1, pages 1241-1275 and 1276-1301, respectively, and references cited therein.

[0384] II.C.11. Ribosomal Targeting

[0385] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in parental cell cytoplasm prior to minicell formation and/or the segregated minicell cytoplasm. By way of non-limiting example, these techniques may include modifications of endogenous and/or exogenous ribosomal components such that ribosomes enter the minicell segregates with higher efficiency. By way of non-limiting example, these techniques may include increasing the copy number of ribosomal binding sites on plasmid or like structure to recruit more ribosomal components or increase the synthesis of ribosomal subunits prior to segregation (Mawn et al., Depletion of free 30S ribosomal subunits in *Escherichia coli* by expression of RNA containing Shine-Dalgarno-like sequences, J. Bacteriol. 184:494-502, 2002). This construct may also include the use of plasmid expressed translation initiation factors to assist ribosomal segregation (Celano et al., Interaction of *Escherichia coli* translation-initiation factor IF-i with ribosomes, Eur. J. Biochem. 178:351-355 1988). See also Hoopes and McClure, "Strategies in Regulation of Transcription Initiation," Chapter 75 in: *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*, Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1987, Volume 2, pages 1231-1240, and references cited therein.

[0386] II.C.12. Proteases

[0387] Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in minicells. By way of non-limiting example, these techniques may include utilization and/or modification of endogenous and/or exogenous proteases. Such manipulations may result in increased or decreased production, and/or changes in the intramolecular and intermolecular functions, of a protein in a minicell or its parent cell; in the latter instance, the protein may be one that is desirably retained in segregated minicells.

[0388] The production or activity of a desired protein gene product may be increased by decreasing the level and/or activity of a protease that acts upon the desired protein. The production or activity of a desired protein gene product may be increased by increasing the level and/or activity of a protease that acts upon a protein that inhibits the production or function of the desired protein.

[0389] The production or activity of a desired nucleic acid gene product may be increased by increasing the level and/or

activity of a protease that acts upon a protein that that inhibits the production or function of the nucleic acid gene product. The production or activity of a desired nucleic acid gene product may be increased by decreasing the level and/or activity of a protease that acts upon a protein that stimulates or enhances the production or function of the desired nucleic acid gene product.

[0390] As one example, manipulation of the *alpA* gene or gene product from *E. coli* (Kirby J. E., and J. E. Trempy, and S. Gottesman. Excision of a P4-like cryptic prophage leads to Alp protease expression in *Escherichia coli*. 1994. J Bacteriol. 176:2068-2081), or homologs of this gene or gene product found in other members of the Prokaryotes, Eukaryotes or Archaeobacteria, may be employed to increase the efficiency of gene expression and protein production in parent cells prior to minicell formation and/or segregated minicells postpartum.

[0391] In addition to *alpA*, other exemplary *E. coli* genes and gene products include the *clpA* gene and gene product from *E. coli* (Katayama Y., and S. Gottesman, and J. Pumphrey, and S. Rudikoff, and W. P. Clark, and M. R. Maurizi. The two-component, ATP-dependent Clp protease of *Escherichia coli*. Purification, cloning, and mutational analysis of the ATP-binding component. 1988. J. Biol. Chem. 263:15226-15236); the *clpB* gene product from *E. coli* (Kitagawa M., and C. Wada, and S. Yoshioka, and T. Yura. Expression of ClpB, an analog of the ATP-dependent protease regulatory subunit in *Escherichia coli*, is controlled by a heat shock sigma factor ( $\sigma$  32). J Bacteriol. 173:4247-4253); the *clpC* gene product from *E. coli* (Msadek T., and F. Kunst, and G. Rapoport. MecB of *Bacillus subtilis*, a member of the ClpC ATPase family, is a pleiotropic regulator controlling competence gene expression and growth at high temperature. 1994. Proc Natl Acad Sci USA 91:5788-5792); the *clpP* gene product from *E. coli* (Maurizi M. R., and W. P. Clark, and Y. Katayama, and S. Rudikoff, and J. Pumphrey, and B. Bowers, and S. Gottesman. Sequence and structure of ClpP, the proteolytic component of the ATP-dependent Clp protease of *Escherichia coli*. 1990. J Biol Chem. 265:12536-12545); the *clpX* gene product from *E. coli* (Gottesman S., and W. P. Clark, and V. de Crecy-Lagard, and M. R. Maurizi. ClpX, an alternative subunit for the ATP-dependent Clp protease of *Escherichia coli*. Sequence and in vivo activities. 1993. J. Biol. Chem. 268:22618-22626); the *clpY* gene product from *E. coli* (Missiakas D., and F. Schwager, J. M. Betton, and C. Georgopoulos, S. Raina. Identification and characterization of HsIV HsIU (ClpQ ClpY) proteins involved in overall proteolysis of misfolded proteins in *Escherichia coli*. 1996. EMBO J. 15:6899-6909); the *dcp* gene product from *E. coli* (Becker S., and Plapp R. Location of the *dcp* gene on the physical map of *Escherichia coli*. 1992. J Bacteriol. 174:1698-1699); the *degP* (*htrA*) gene product from *E. coli* (Lipinska B., and M. Zylicz, and C. Georgopoulos. The HtrA (DegP) protein, essential for *Escherichia coli* survival at high temperatures, is an endopeptidase. 1990. J Bacteriol. 172:1791-1797); the *ggt* gene product from *E. coli* (Finidori J., and Y. Laperche, and R. Haguenaer-Tsapis, and R. Barouki, and G. Guellaen, and J. Hanoune. In vitro biosynthesis and membrane insertion of gamma-glutamyl transpeptidase. 1984. J. Biol. Chem. 259:4687-4690); the *hfl* gene product from *E. coli* (Cheng H. H., and H. Echols. A class of *Escherichia coli* proteins controlled by the *hflA* locus. 1987. J. Mol. Biol. 196:737-740); the *hflB* gene

product from *E. coli* (Banuett F., and M. A. Hoyt, and L. McFarlane, and H. Echols, and I. Herskowitz. *HflB*, a new *Escherichia coli* locus regulating lysogeny and the level of bacteriophage lambda c11 protein. 1986. J. Mol. Biol. 187:213-224); the *hflC* gene product from *E. coli* (Noble J. A., and M. A. Innis, and E. V. Koonin, and K. E. Rudd, and F. Banuett, and I. Herskowitz, The *Escherichia coli* *hflA* locus encodes a putative GTP-binding protein and two membrane proteins, one of which contains a protease-like domain. 1993. Proc Natl Acad Sci USA. 90:10866-10870); the *hflK* gene product from *E. coli* (Id.); the *hflX* gene product from *E. coli* (Noble J. A., and M. A. Innis, and E. V. Koonin, and K. E. Rudd, and F. Banuett, and I. Hertzskowitz. The *Escherichia coli* *hflA* locus encodes a putative GTP-binding protein and two membrane proteins, one of which contains a protease-like domain. 1993. Proc Natl Acad Sci USA. 90:10866-10870); the *hopD* gene product from *E. coli* (Whitchurch C. B., and J. S. Mattick. *Escherichia coli* contains a set of genes homologous to those involved in protein secretion, DNA uptake and the assembly of type-4 fimbriae in other bacteria. 1994. Gene. 150:9-15); the *htrA* gene product from *E. coli* (Lipinska B., and S. Sharma, and C. Georgopoulos. Sequence analysis and regulation of the *htrA* gene of *Escherichia coli*: a sigma 32-independent mechanism of heat-inducible transcription. 1988. Nucleic Acids Res. 16:10053-10067); the *hycI* gene product from *E. coli* (Rossmann R., and T. Maier, and F. Lottspeich, and A. Bock. Characterisation of a protease from *Escherichia coli* involved in hydrogenase maturation. 1995. Eur J Biochem. 227:545-550); the *iap* gene product from *E. coli* (Nakata A., and M. Yamaguchi, and K. Isutani, and M. Amemura. *Escherichia coli* mutants deficient in the production of alkaline, phosphatase isozymes. 1978. J Bacteriol. 134:287-294); the *lep* gene product from *E. coli* (Silver P., and W. Wickner. Genetic mapping of the *Escherichia coli* leader (signal) peptidase gene (*lep*): a new approach for determining the map position of a cloned gene. 1983. J Bacteriol. 54:659-572); the *Ion* gene product from *E. coli* (Donch J., and J. Greenberg. Genetic analysis of *Ion* mutants of strain K-12 of *Escherichia coli*. 1968. Mol Gen Genet. 103:105-115); the *lsp* gene product from *E. coli* (Regue M., and J. Remenick, and M. tokunaga, and G. A. Mackie, and H. C. Wu. Mapping of the lipoprotein signal peptidase gene (*Isp*). 1984. J Bacteriol. 194:632-635); the *ompT* gene product from *E. coli* (Akiyama Y., and K. SecY protein, a membrane-embedded secretion factor of *E. coli*, is cleaved by the *ompT* protease in vitro. 1990. Biochem Biophys Res Commun. 167:711-715); the *opdA* gene product from *E. coli* (Conllin C. A., and C. G. Miller. Location of the *prlC* (*opdA*) gene on the physical map of *Escherichia coli*. 1993. J Bacteriol. 175:5731-5732); the *orfX* gene product from *E. coli* (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umberger (eds.). *Escherichia coli* and *Salmonella typhimurium: Cellular and Molecular Biology*, 2<sup>nd</sup> ed. American Society for Microbiology, Washington D.C.); the *pepA* gene product from *E. coli* (Stirling C. J., and S. D. Colloms, and J. F. Collins, and G. Szatmari, and D. J. Sherratt. *XerB*, an *Escherichia coli* gene required for plasmid ColE1 site-specific recombination, is identical to *pepA*, encoding aminopeptidaseA, a protein with substantial similarity to bovine lens leucine aminopeptidase. 1989. EMBO J. 8:1623-1627);

- the pepD gene product from *E. coli* (Henrich B., and U. Schroeder, and R. W. Frank, and R. Plapp. Accurate mapping of the *Escherichia coli* pepD gene by sequence analysis of its 5' flanking region. 1989. *Mol Gen Genet.* 215:369-373); the pepE gene product from *E. coli* (Conlin C. A., and T. M. Knox, and C. G. Miller. Cloning and physical map position of an alpha-aspartyl depeptidase gene, pepE, from *Escherichia coli*. 1994. *J Bacteriol.* 176:1552-1553); the pepN gene product from *E. coli* (Miller C. G., and G. Schwartz. Peptidase-deficient mutants of *Escherichia coli*. 1978. *J Bacteriol.* 135:603-611); the pepP gene product from *E. coli* (Id.); the pepQ gene product from *E. coli* (Id.); the pepT gene product from *E. coli* (Miller G. G., and G. Schwartz. Peptidase-deficient mutants of *Escherichia coli*. 1978. *J Bacteriol.* 135:603-611); the pilD gene product from *E. coli* (Francetic O., and S. Lory, and A. P. Pugsley. A second prepilin peptidase gene in *Escherichia coli* K-12. 1998. *Mol Microbiol.* 27:763-775); the pinA gene product from *E. coli* (Hilliard J. J., and L. D. Simon, and L. Van Melderen, and M. R. Maurizi. PinA inhibits ATP hydrolysis and energy-dependent protein degradation by Lon protease. 1998. *J. Biol. Chem.* 273:524-527); the prc(tsp) gene product from *E. coli* (Nagasawa H., and Y. Sakagami, and A. Suzuki, and H. Suzuki, and H. Hara, and Y. Hirota. Determination of the cleavage site involved in C-terminal processing of penicillin-binding protein 3 of *Escherichia coli*. 1989. *J Bacteriol.* 171:5890-5893); the prlC gene product from *E. coli* (Jiang X., and M. Zhang, and Y. Ding, and J. Yao, and H. Chen, and D. Zhu, and M. Muramatu. *Escherichia coli* prlC gene encodes a trypsin-like proteinase regulating the cell cycle. 1998. *J Biochem (Tokyo)* 128:980-985); the protease V gene product from *E. coli* (Berlyn, M. K. B. et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9, In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2<sup>nd</sup> ed. American Society for Microbiology, Washington, D.C.); the protease VI gene product from *E. coli* (Id.); the protease In gene product from *E. coli* (Id.); the protease Fa gene product from *E. coli* or homologues (Id.); the protease Mi gene product from *E. coli* (Id.); the protease So gene product from *E. coli* (Id.); the ptrA gene product from *E. coli* (Id.); the ptrB gene product from *E. coli* (Id.); the sypB gene product from *E. coli* (Barends S., and A. W. Karzai, and R. T. Sauer, and J. Wower, and B. Kraal. Simultaneous a functional binding of SmpB and EF-Tu-TP to the analyl acceptor arm of tmRNA. 2001. *J. Mol. Biol.* 314:9-21); the sohB gene product from *E. coli* (Baird L., and B. Lipinska, and S. Raina, and C. Georgopoulos. Identification of the *Escherichia coli* sohB gene, a multicopy suppressor of the HtrA (DegP) null phenotype. 1991. *J Bacteriol.* 173-5763-5770); the sspA gene product from *E. coli* (Ichihara S., and T. Suzuki, and M. Suzuki, and C. Mizushima. Molecular cloning and sequencing of the sppA gene and characterization of the encoded protease IV, a signal peptide peptidase of *Escherichia coli*. 1986. *J. Biol. Chem.* 261:9405-9411); the tesA gene product from *E. coli* (Cho H., and J. E. Cronan Jr. *Escherichia coli* thioesterase I, molecular cloning and sequencing of the structural gene and identification as a periplasmic enzyme. 1993. *J. Biol. Chem.* 268:9238-9245); the tufA gene product from *E. coli* (Ang., and J. S. Lee, and J. D. Friesen. Evidence for an internal promoter preceding tufA in the str operon of *Escherichia coli*. *J Bacteriol.* 149:548-553); the tufB gene product from *E. coli* (Mihajima A., and M. Shibuya, and Y. Kaziro. Construction and characterization of the two hybrid Col1E1 plasmids carrying *Escherichia coli* tufB gene. 1979. *FEBS Lett.* 102:207-210); the ybau gene product from *E. coli* (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2<sup>nd</sup> ed. American Society for Microbiology, Washington, D.C.); the ssrA gene (tmRNA, 10Sa RNA) product from *E. coli* (Oh B. K., and A. K. Chauhan, and K. Isono, and D. Apirion. Location of a gene (ssrA) for a small, stable RNA 910Sa RNA) in the *Escherichia coli* chromosome. 1990. *J Bacteriol.* 172:4708-4709); and the ssrB gene from *E. coli* (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger 9eds.). *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2<sup>nd</sup> ed. American Society for Microbiology, Washington, D.C.).
- [0392]** II.C.13. Chaperones
- [0393]** Included in the design of the invention are techniques that increase the efficiency of gene expression and functional protein production in minicells. By way of non-limiting example, these techniques may include modification of chaperones and chaperonins, i.e., endogenous and/or exogenous protein components that monitor the unfolded state of translated proteins allowing proper folding and/or secretion, membrane insertion, or soluble multimeric assembly of expressed proteins in the parental cell prior to minicell formation and/or the segregated minicell cytoplasm, membrane, periplasm, and/or extracellular environment. See Gottesman et al., Protein folding and unfolding by *Escherichia coli* chaperones and chaperonins, *Current Op. Microbiol.* 3:197-202, 2000; and Mayhew et al., "Molecular Chaperone Proteins," Chapter 61 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2nd Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 1, pages 922-937, and references cited therein.
- [0394]** These applications may, but are not limited to increased or decreased chaperone production, increased or decreased intramolecular activity of a chaperone, increased or decreased physiological function of a chaperone, or deletion, substitution, inversion, translocation or insertion into, or mutation of, a gene encoding a chaperone. By way of non-limiting example, increased production of a chaperone may occur through increased chaperone gene dosage (increased copy number of a given gene under the control of the native or artificial promoter where this gene may be on a plasmid or in more than one copy on the chromosome), modification of the native regulatory elements, including, but not limited to the promoter or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/enhancers, or relevant antisense nucleic acid or nucleic acid analog, cloning on a plasmid under the control of the native or artificial promoter, and increased or decreased production of native or artificial promoter regulatory element(s) controlling production of

the chaperone gene or gene product. By way of non-limiting example, decreased production of a chaperone may occur through modification of the native regulatory elements, including, but not limited to the promotor or operator region(s) of DNA, or ribosomal binding sites on RNA, relevant repressors/silencers, relevant activators/enhancers, or relevant antisense nucleic acid or nucleic acid analog, through cloning on a plasmid under the control of the native regulatory region containing mutations or an artificial promotor, either or both of which resulting in decreased chaperone production, and through increased or decreased production of native or artificial promotor regulatory element(s) controlling production of the chaperone gene or gene product. By definition, intramolecular activity refers to the enzymatic function, structure-dependent function, e.g. the capacity of chaperone to interact in a protein-protein, protein-nucleic acid, or protein-lipid complex, and/or carrier function, e.g. the capacity to bind, either covalently or non-covalently small organic or inorganic molecules, protein(s), carbohydrate(s), fatty acid(s), lipid(s), and nucleic acid(s). By way of non-limiting example, alteration of intramolecular activity may be accomplished by mutation of the chaperone gene, in vivo or in vitro chemical modification of Chaperone, inhibitor molecules against the function of chaperone, e.g. competitive, non-competitive, or uncompetitive enzymatic inhibitors, inhibitors that prevent protein-protein, protein-nucleic acid, or protein-lipid interactions, e.g. expression or introduction of dominant-negative or dominant-positive chaperone or other protein fragment(s), or other carbohydrate(s), fatty acid(s), lipid(s), and nucleic acid(s) that may act directly or allosterically upon Chaperone, and/or modification of protein, carbohydrate, fatty acid, lipid, or nucleic acid moieties that modify the chaperone gene or gene product to create the functional protein. By definition, physiological function refers to the effects resulting from an intramolecular interaction between Chaperone and other protein, carbohydrate, fatty acid, lipid, nucleic acid, or other molecule(s) in or on the cell or the action of a product or products resulting from such an interaction. By way of non-limiting example, physiological function may be the act or result of intermolecular phosphorylation, biotinylation, methylation, acylation, glycosylation, and/or other signaling event; this function may be the result of a protein-protein, protein-nucleic acid, or protein-lipid interaction resulting in a functional moiety; this function may be to interact with the membrane to recruit other molecules to this compartment of the cell; this function may be to regulate the transcription and/or translation of chaperone, other protein, or nucleic acid; and this function may be to stimulate the function of another process that is not yet described or understood.

[0395] By way of non-limiting example, chaperone genes may be any of the *E. coli* genes listed below, as well as any homologs thereof from prokaryotes, eukaryotes, archaeobacteria, or organelles (mitochondria, chloroplasts, plastids, etc.). Exemplary *E. coli* genes encoding chaperones include, by way of non-limiting example, the *cbpA* gene (Shiozawa T., and C. Ueguchi, and T. Mizuno. The *rpoD* gene functions as a multicopy suppressor for mutations in the chaperones, *CbpA*, *DnaJ* and *DnaK*, in *Escherichia coli*. 1996 FEMS Microbiol Lett. 138:245-250); the *clpB* gene (Squires C. L., and S. Pedersen, and B. M. Ross, and C. Squires. *ClpB* is the *Escherichia coli* heat shock protein F84. 1. 1991. J Bacteriol. 173:4254-4262); the *draK* gene (Kroczyńska B., and S.

Y. Blond. Cloning and characterization of a new soluble murine J-domain protein that stimulates BiP, Hsc70 and DnaK ATPase activity with different efficiencies. 2001. Gene. 273:267-274); the *dnaJ* gene (Kedzierska S., and E. Matuszewska. The effect of co-overproduction of DnaK/DnaJ/GrpE and ClpB proteins on the removal of heat-aggregated proteins from *Escherichia coli* Delta *clpB* mutant cells—new insight into the role of Hsp70 in a functional cooperation with Hsp100. 2001. FEMS Microbiol Lett. 204:355-360); the *ecpD* gene (Raina S., and D. Missiakas, and L. Baird, and S. Kumar, and C. Georgopoulos. Identification and transcriptional analysis of the *Escherichia coli* *htrE* operon which is homologous to *pap* and related pilin operons. 1993. J Bacteriol. 175:5009-5021); the *ffh* gene (Muller, M., et al. 1002. Protein traffic in bacteria: multiple routes from the ribosome to and across the membrane. Prog. Nucleic Acid Res. Mol. Biol. 66:107-157); 4.5S RNA (Muller, M., et al. 1002. Protein traffic in bacteria: multiple routes from the ribosome to and across the membrane. Prog. Nucleic Acid Res. Mol. Biol. 66:107-157); the *FtsY* gene (Muller, M., et al. 1002. Protein traffic in bacteria: multiple routes from the ribosome to and across the membrane. Prog. Nucleic Acid Res. Mol. Biol. 66:107-157); the *fimC* gene (Klemm P., and B. J. Jorgensen, and 1. van Die, and H. de Ree, and H. Bergmans. The *fim* genes responsible for synthesis of type 1 fimbriae in *Escherichia coli*, cloning and genetic organization. 1985. Mol Gen Genet. 199:410-414); the *groE* gene (Burton Z. F., and D. Eisenberg. A procedure for rapid isolation of both *groE* protein and glutamine synthetase from *E. coli*. 1980. Arch Biochem Biophys. 205:478-488); the *groL* gene (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli* and *Salmonella typhimurium: cellular and molecular biology*, 2nd ed. American Society for Microbiology, Washington D.C.); the *groS* gene (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli* and *Salmonella typhimurium: cellular and molecular biology*, 2nd ed. American Society for Microbiology, Washington D.C.); the *hptG* gene (Berlyn, M. K. B., et al. 1996. Linkage map of *Escherichia coli* K-12, Edition 9. In F. C. Neidhardt, R. Curtiss, J. L. Ingraham, E. C. C. Lin, K. B. Low, B. Magasanik, W. S. Reznikoff, M. Riley, M. Schaechter, and H. E. Umbarger (eds.). *Escherichia coli* and *Salmonella typhimurium: cellular and molecular biology*, 2nd ed. American Society for Microbiology, Washington D.C.); the *hscA* gene (Takahashi Y., and M. Nakamura. Functional assignment of the ORF2-iscS-iscU-iscA-hscB-hscA-*fdx*-ORF3 gene cluster involved in the assembly of Fe-S clusters in *Escherichia coli*. 1999. J Biochem (Tokyo). 126:917-926); the *ibpA* gene (Lund P. A. Microbial molecular chaperones. 2001. Adv Microb Physiol. 44:93-140); the *papJ* gene (Tennent, J. M., et al. 1990. Integrity of *Escherichia coli* P pili during biogenesis: properties and role of PapJ. Mol. Microbiol. 4:747-758); the *secB* gene (Lecker, S., et al. 1989. Three pure chaperone proteins of *Escherichia coli*—SecB, trigger factor and GroEL—form soluble complexes with precursor proteins in vitro. EMBO J. 8:2703-2709); and the *tig* gene (Lecker, S., et al. 1989. Three pure chaperone



proteins of *Escherichia coli*—SecB, trigger factor and GroEL—form soluble complexes with precursor proteins in vitro. EMBO J. 8:2703-2709); the secE gene (Muller, M., et al. 1002. Protein traffic in bacteria: multiple routes from the ribosome to and across the membrane. Prog. Nucleic Acid Res. Mol. Biol. 66:107-157); and the secY gene (Muller, M., et al. 1002. Protein traffic in bacteria: multiple routes from the ribosome to and across the membrane. Prog. Nucleic Acid Res. Mol. Biol. 66:107-157).

**[0396]** II.C.14. Export Apparatus and Membrane Targeting

**[0397]** Included in the design of the invention are techniques that increase the efficiency of gene expression and protein production in parental cells prior to minicell formation and/or in the segregated minicells. By way of non-limiting example, these techniques may include construction of chimeric proteins including, but not limited to, coupling the expressed protein of interest with native Eubacterial, Eukaryotic, Archeobacterial or organellar leader sequences to drive membrane insertion or secretion of the protein of interest to the periplasm or extracellular environment. In addition to using native leader sequences, these minicell expression constructs may also include proteolytic cleavage sites to remove the leader sequence following insertion into the membrane or secretion. These proteolytic cleavage sites may be native to the organism from which the minicell is derived or non-native. In the latter example, also included in the system are the non-native protease that recognizes the non-native proteolytic cleavage site.

**[0398]** Non-limiting examples of these leader sequences may be the leader from the STII protein (Voss, T., et al. 1994. Periplasmic expression of human interferon-alpha 2c in *Escherichia coli* results in a correctly folded molecule. Biochem. J. 298:719-725), maltose binding protein (malE) (Ito, K. 1982. Purification of the precursor form of maltose-binding protein, a periplasmic protein of *Escherichia coli*. J. Biol. Chem. 257:9895-9897), phoA (Jobling, M. G., et al. 1997. Construction and characterization of versatile cloning vectors for efficient delivery of native foreign proteins to the periplasm of *Escherichia coli*. Plasmid. 38:158-173), lamb (Wong, E. Y., et al. 1988. Expression of secreted insulin-like growth factor-i in *Escherichia coli*. Gene. 68:193-203), ompA (Loe, T., et al. 2002. Using secretion to solve a solubility problem: high-yield expression in *Escherichia coli* and purification of the bacterial glycoamidase PNGase F. Protein Expr. Purif. 24:90-98), or pelB (Molloy, P. E., et al. 1998. Production of soluble single-chain T-cell receptor fragments in *Escherichia coli* trxB mutants. Mol. Immunol. 35:73-81).

**[0399]** In addition to these leader sequences, mutations in the cellular export machinery may be employed to increase the promiscuity of export to display or export sequences with non-optimized leader sequences. Non-limiting examples of genes that may be altered to increase export promiscuity are mutations in secY (prlA4) (Derman, A. I., et al. 1993. A signal sequence is not required for protein export in prlA mutants of *Escherichia coli*. EMBO J. 12:879-888), and secE (Harris, C. R., and T. J. Silhavy. 1999. Mapping an interface of SecY (PrlA) and SecE (PrlG) by using synthetic phenotypes and in vivo cross-linking. J. Bacteriol. 181:3438-3444).

**[0400]** II.C.15. Increasing Stability and Solubility

**[0401]** Included in the design of the invention are techniques that increase the efficiency of gene expression and

protein production in parental cells prior to minicell formation and/or in the segregated minicells. By way of non-limiting example, these techniques may include construction of chimeric/fusion proteins including, but not limited to, coupling the expressed protein of interest with native Eubacterial, Eukaryotic, Archeobacterial or organellar solubilizing sequences. As used herein, "solubilizing sequences" are complete or truncated amino acid sequences that increase the solubility of the expressed membrane protein of interest. This increased solubility may be used to increase the lifetime of the soluble state until proper membrane insertion may take place. By way of non-limiting example, these soluble chimeric fusion proteins may be ubiquitin (Power, R. F., et al. 1990. High level expression of a truncated chicken progesterone receptor in *Escherichia coli*. J. Biol. Chem. 265:1419-1424), thioredoxin (LaVallie, E. R., et al. 1993. A thioredoxin gene fusion expression system that circumvents inclusion body formation in the *E. coli* cytoplasm. Biotechnology (N.Y.) 11:187-193; Kapust, R. B., and D. S. Waugh. 1999. *Escherichia coli* maltose-binding protein is uncommonly effective at promoting the solubility of polypeptides to which it is fused. Protein Sci. 8:1668-1674), the dsbA gene product (Winter, J., et al. 2001. Increased production of human proinsulin in the periplasmic space of *Escherichia coli* by fusion to DsbA. J. Biotechnol. 84:175-185), the SPG protein (Murphy, J. P., et al. 1992. Amplified expression and large-scale purification of protein G'. Bioseparation 3:63-71), the malE gene product (maltose-binding protein) (Hampe, W., et al. 2000. Engineering of a proteolytically stable human beta 2-adrenergic receptor/maltose-binding protein fusion and production of the chimeric protein in *Escherichia coli* and baculovirus-infected insect cells. J. Biotechnol. 77:219-234; Kapust et al., *Escherichia coli* maltose-binding protein is uncommonly effective at promoting the solubility of polypeptides to which it is fused, Protein Sci. 8:1668-1674, 1999), glutathione-s-transferase (GST); and/or nuclease A (Meeker et al., A fusion protein between serum amyloid A and staphylococcal nuclease—synthesis, purification, and structural studies, Proteins 30:381-387, 1998). In addition to these proteins, Staphylococcal protein A, beta-galactosidase, S-peptide, myosin heavy chain, dihydrofolate reductase, T4 p55, growth hormone N terminus, *E. coli* Hemolysin A, bacteriophage lambda cII protein, TrpE, and TrpLE proteins may also be used as fusion proteins to increase protein expression and/or solubility (Makrides, Strategies for Achieving High-Level Expression of Genes in *Escherichia coli*, Microbiol. Rev. 60:512-538).

**[0402]** III. Preparation of Minicells

**[0403]** III.A. Parent Cell Mutations

**[0404]** Although it has been reported that relatively few molecules of endogenous RNA polymerase segregate into minicells (Shepherd et al., Cytoplasmic RNA Polymerase in *Escherichia coli*, J Bacteriol 183:2527-34, 2001), other reports and results indicate that many RNA Polymerase molecules follow plasmids into minicells (Funnell and Gagnier, Partition of P1 plasmids in *Escherichia coli* mukB chromosomal partition mutants, J Bacteriol 177:2381-6, 1995). In any event, applicants have discovered that the introduction of an exogenous RNA polymerase to minicell-producing cells enhances expression of episomal elements in minicells. Such enhanced expression may allow for the



successful expression of proteins in minicells, wherein such proteins are expressed poorly or not at all in unmodified minicells. In order to maximize the amount of RNA transcription from episomal elements in minicells, minicell-producing cell lines that express an RNA polymerase specific for certain episomal expression elements may be used. An example of an *E. coli* strain of this type, designated MC-T7, was created and used as is described in the Examples. Those skilled in the art will be able to make and use equivalent strains based on the present disclosure and their knowledge of the art.

[0405] Minicell-producing cells may comprise mutations that augment preparative steps. For example, lipopolysaccharide (LPS) synthesis in *E. coli* includes the lipid A biosynthetic pathway. Four of the genes in this pathway have now been identified and sequenced, and three of them are located in a complex operon that also contains genes involved in DNA and phospholipid synthesis. The *rfa* gene cluster, which contains many of the genes for LPS core synthesis, includes at least 17 genes. The *rfb* gene cluster encodes protein involved in O-antigen synthesis, and *rfb* genes have been sequenced from a number of serotypes and exhibit the genetic polymorphism anticipated on the basis of the chemical complexity of the O antigens. See Schnaitman and Klena, Genetics of lipopolysaccharide biosynthesis in enteric bacteria, *Microbiol. Rev.* 57:655-82, 1993. When present, alone, or in combination, the *rfb* and *oms* mutations cause alterations in the eubacterial membrane that make it more sensitive to lysozyme and other agents used to process minicells. Similarly, the *rfa* (Chen, L., and W. G. Coleman Jr. 1993. Cloning and characterization of the *Escherichia coli* K-12 *rfa-2* (*rfaC*) gene, a gene required for lipopolysaccharide inner core synthesis. *J. Bacteriol.* 175:2534-2540), *IpcA* (Brooke, J. S., and M. A. Valvano. 1996. Biosynthesis of inner core lipopolysaccharide in enteric bacteria identification and characterization of a conserved phosphoheptose isomerase. *J. Biol. Chem.* 271:3608-3614), and *IpcB* (Kadrman, J. L., et al. 1998. Cloning and overexpression of glycosyltransferases that generate the lipopolysaccharide core of *Rhizobium leguminosarum*. *J. Biol. Chem.* 273:26432-26440) mutations, when present alone or in combination, cause alterations in lipopolysaccharides in the outer membrane causing cells to be more sensitive to lysozyme and agents used to process minicells. In addition, such mutations can be used to reduce the potential antigenicity and/or toxicity of minicells.

#### [0406] III.B. Culturing Conditions

[0407] Included in the design of the invention are the conditions to grow parental cells from which minicells will be produced. Non-limiting examples herein are drawn to conditions for growing *E. coli* parental cells to produce minicells derived from *E. coli* parental cells. Non-limiting examples for growth media may include rich media, e.g. Luria broth (LB), defined minimal media, e.g. M63 salts with defined carbon, nitrogen, phosphate, magnesium, and sulfate levels, and complex minimal media, e.g. defined minimal media with casamino acid supplement. This growth may be performed in culture tubes, shake flasks (using a standard air incubator, or modified bioreactor shake flask attachment), or bioreactor. Growth of parental cells may include supplemented additions to assist regulation of expression constructs listed in the sections above. These supplements may include, but are not limited to dextrose,

phosphate, inorganic salts, ribonucleic acids, deoxyribonucleic acids, buffering agents, thiamine, or other chemical that stimulates growth, stabilizes growth, serves as an osmoprotectant, regulates gene expression, and/or applies selective pressure to mutation, and/or marker selection. These mutations may include an amino acid or nucleotide auxotrophy, while the selectable marker may include transposable elements, plasmids, bacteriophage, and/or auxotrophic or antibiotic resistance marker. Growth conditions may also require temperature adjustments, carbon alternations, and/or oxygen-level modifications to stimulate temperature sensitive mutations found in designed gene products for a given desired phenotype and optimize culture conditions.

[0408] By way of non-limiting example, production of minicells and protein production may occur by using either of two general approaches or any combination of each. First, minicells may be formed, purified, and then contained expression elements may be stimulated to produce their encoded gene products. Second, parental cells, from which the minicells are derived, may be stimulated to express the protein of interest and segregate minicells simultaneously. Finally, any timing variable of minicell formation and protein production may be used to optimize protein and minicell production to best serve the desired application. The two general approaches are shown in the sections below.

#### [0409] III.C. Manipulation of Genetic Expression in Minicell Production

[0410] Included in the design of the invention are methods that increase the efficiency, rate and/or level of gene expression and protein production in parent cells and/or minicells. Such methods include, but are not limited to, the following.

[0411] By way of non-limiting example, parental cells are grown overnight in the appropriate media. From this culture, the cells are subcultured into the same media and monitored for growth. At the appropriate cell density, the cultures are induced for minicell production using any of the switching mechanisms discussed in section II.B. regulating any construct discussed in section II.A. Non-limiting examples of this minicell-inducing switching mechanism may be the *ileR* gene product regulating the production of the *hns* minicell-inducing gene product or the *melR* gene product regulating the production of the *minB* minicell-inducing gene product. Following minicell induction, the culture is allowed to continue growth until the desired concentration of minicells is obtained. At this point, the minicells are separated from the parental cells as described in section II.E. Purified minicells are induced for protein production by triggering the genetic switching mechanism that segregated into the minicell upon separation from the parental cell. By way of non-limiting example, this genetic switching mechanism may be any of those discussed in section I.B. regulating the production of any protein of interest. Furthermore, at this point or during the production of minicells the peripheral gene expression, production, and assembly machinery discussed in section II.C. may be triggered to assist in this process. By way of non-limiting example, the *groEL* complex may be triggered using the temperature sensitive *lambda cI* inducible system from a co-segregant plasmid to assist in the proper assembly of the expressed protein of interest.

#### [0412] III.D. Separation of Minicells from Parent Cells

[0413] A variety of methods are used to separate minicells from parent cells (i.e., the cells from which the minicells are

produced) in a mixture of parent cells and minicells. In general, such methods are physical, biochemical and genetic, and can be used in combination.

**[0414]** III.D.1. Physical Separation of Minicells from Parent Cells

**[0415]** By way of non-limiting example, minicells are separated from parent cells glass-fiber filtration (Christen et al., *Gene* 23:195-198, 1983), and differential and zonal centrifugation (Barker et al., *J. Gen. Microbiol.* 111:387-396, 1979), size-exclusion chromatography, e.g. gel-filtration, differential sonication (Reeve, J. N., and N. H. Mendelson. 1973. Pronase digestion of amino acid binding components on the surface of *Bacillus subtilis* cells and minicells. *Biochem. Biophys. Res. Commun.* 53:1325-1330), and UV-irradiation (Tankersley, W. G., and J. M. Woodward. 1973. Induction and isolation of non-replicative minicells of *Salmonella typhimurium* and their use as immunogens in mice. *Bacteriol. Proc.* 97).

**[0416]** Some techniques involve different centrifugation techniques, e.g., differential and zonal centrifugation. By way of non-limiting example, minicells may be purified by the double sucrose gradient purification technique described by Frazer and Curtiss, *Curr. Topics Microbiol. Immunol.* 69:1-84, 1975. The first centrifugation involves differential centrifugation, which separates parent cells from minicells based on differences in size and/or density. The percent of sucrose in the gradient (graduated from about 5 to about 20%), Ficoll or glycerol is designed to allow only parent cells to pass through the gradient.

**[0417]** The supernatant, which is enriched for minicells, is then separated from the pellet and is spun at a much higher rate (e.g.,  $\geq 11,000$  g). This pellets the minicells and any parent cells that did not pellet out in the first spin. The pellet is then resuspended and layered on a sucrose gradient.

**[0418]** The band containing minicells is collected, pelleted by centrifugation, and loaded on another gradient. This procedure is repeated until the minicell preparation is essentially depleted of parent cells, or has a concentration of parent cells that is low enough so as to not interfere with a chosen minicell application or activity. By way of non-limiting example, buffers and media used in these gradient and resuspension steps may be LB, defined minimal media, e.g. M63 salts with defined carbon, nitrogen, phosphate, magnesium, and sulfate levels, complex minimal media, e.g. defined minimal media with casamino acid supplement, and/or other buffer or media that serves as an osmo-protectant, stabilizing agent, and/or energy source, or may contain agents that limit the growth of contaminating parental cells, e.g. azide, antibiotic, or lack an auxotrophic supplemental requirement, e.g. thiamine.

**[0419]** Other physical methods may also be used to remove parent cells from minicell preparations. By way of non-limiting example, mixtures of parent cells and minicells are frozen to  $-20^{\circ}$  C. and then thawed slowly (Frazer and Curtiss, *Curr. Topics Microbiol. Immunol.* 69:1-84, 1975).

**[0420]** III.D.2. Biochemical Separation of Minicells from Parent Cells

**[0421]** Contaminating parental cells may be eliminated from minicell preparations by incubation in the presence of an agent, or under a set of conditions, that selectively kills

dividing cells. Because minicells can neither grow nor divide, they are resistant to such treatments.

**[0422]** Examples of biochemical conditions that prevent or kill dividing parental cells is treatment with an antibacterial agent, such as penicillin or derivatives of penicillin. Penicillin has two potential effects. First, penicillin prevent cell wall formation and leads to lysis of dividing cells. Second, prior to lysis dividing cells form filaments that may assist in the physical separation steps described in section III.E.1. In addition to penicillin and its derivatives, other agents may be used to prevent division of parental cells. Such agents may include azide. Azide is a reversible inhibitor of electron transport, and thus prevents cell division. As another example, D-cycloserine or phage MS2 lysis protein may also serve as a biochemical approach to eliminate or inhibit-dividing parental cells. (Markiewicz et al., *FEMS Microbiol. Lett.* 70:119-123, 1992). Khachatourians (U.S. Pat. No. 4,311,797) states that it may be desirable to incubate minicell/parent cell mixtures in brain heart infusion broth at  $36^{\circ}$  C. to  $38^{\circ}$  C. prior to the addition of penicillin G and further incubations.

**[0423]** III.D.3. Genetic Separation of Minicells from Parent Cells

**[0424]** Alternatively or additionally, various techniques may be used to selectively kill, preferably lyse, parent cells. For example, although minicells can internally retain M13 phage in the plasmid stage of the M13 life cycle, they are refractory to infection and lysis by M13 phage (Staudenbauer et al., *Mol. Gen. Genet.* 138:203-212, 1975). In contrast, parent cells are infected and lysed by M13 and are thus are selectively removed from a mixture comprising parent cells and minicells. A mixture comprising parent cells and minicells is treated with M13 phage at an M.O.I.=5 (phage:cells). The infection is allowed to continue to a point where  $\geq 50\%$  of the parent cells are lysed, preferably  $\geq 75\%$ , more preferably  $\geq 95\%$  most preferably  $\geq 99\%$ ; and  $\leq 25\%$  of the minicells are lysed or killed, preferably  $<15\%$ , most preferably  $<1\%$ .

**[0425]** As another non-limiting example of a method by which parent cells can be selectively killed, and preferably lysed, a chromosome of a parent cell may include a conditionally lethal gene. The induction of the chromosomal lethal gene will result in the destruction of parent cells, but will not affect minicells as they lack the chromosome harboring the conditionally lethal gene. As one example, a parent cell may contain a chromosomal integrated bacteriophage comprising a conditionally lethal gene. One example of such a bacteriophage is an integrated lambda phage that has a temperature sensitive repressor gene (e.g., lambda c1857). Induction of this phage, which results in the destruction of the parent cells but not of the achromosomal minicells, is achieved by simply raising the temperature of the growth media. A preferred bacteriophage to be used in this method is one that kills and/or lyses the parent cells but does not produce infective particles. One non-limiting example of this type of phage is one that lyses a cell but which has been engineered to as to not produce capsid proteins that are surround and protect phage DNA in infective particles. That is, capsid proteins are required for the production of infective particles.

**[0426]** As another non-limiting example of a method by which parent cells can be selectively killed or lysed, toxic

proteins may be expressed that lead to parental cell lysis. By way of non-limiting example, these inducible constructs may employ a system described in section II.B. to control the expression of a phage holing gene. Holin genes fall with in at least 35 different families with no detectable orthologous relationships (Grundling, A., et al. 2001. Holins kill without warning. Proc. Natl. Acad. Sci. 98:9348-9352) of which each and any may be used to lyse parental cells to improve the purity of minicell preparations.

[0427] Gram negative eubacterial cells and minicells are bounded by an inner membrane, which is surrounded by a cell wall, wherein the cell wall is itself enclosed within an outer membrane. That is, proceeding from the external environment to the cytoplasm of a minicell, a molecule first encounters the outer membrane (OM), then the cell wall and finally, the inner membrane (IM). In different aspects of the invention, it is preferred to disrupt or degrade the OM, cell wall or IM of a eubacterial minicell. Such treatments are used, by way of non-limiting example, in order to increase or decrease the immunogenicity, and/or to alter the permeability characteristics, of a minicell.

[0428] Eubacterial cells and minicells with altered membranes and/or cell walls are called "poroplasts<sup>TM</sup>", "spheroplasts," and "protoplasts." Herein, the terms "spheroplast" and "protoplast" refer to spheroplasts and protoplasts prepared from minicells. In contrast, "cellular spheroplasts" and "cellular protoplasts" refer to spheroplasts and protoplasts prepared from cells. Also, as used herein, the term "minicell" encompasses not only minicells per se but also encompasses poroplasts, spheroplasts and protoplasts.

[0429] In a poroplast, the eubacterial outer membrane (OM) and LPS have been removed. In a spheroplast, portions of a disrupted eubacterial OM and/or disrupted cell wall either may remain associated with the inner membrane of the minicell, but the membrane and cell wall is nonetheless porous because the permeability of the disrupted OM and cell wall has been increased. A membrane is said to be "disrupted" when the membrane's structure has been treated with an agent, or incubated under conditions, that leads to the partial degradation of the membrane, thereby increasing the permeability thereof. In contrast, a membrane that has been "degraded" is essentially, for the applicable intents and purposes, removed. In preferred embodiments, irrespective of the condition of the OM and cell wall, the eubacterial inner membrane is not disrupted, and membrane proteins displayed on the inner membrane are accessible to compounds that are brought into contact with the minicell, poroplast, spheroplast, protoplast or cellular poroplast, as the case may be.

[0430] III.E.2. Poroplasts<sup>TM</sup>

[0431] For various applications poroplasted minicells are capable of preserving the cytoplasmic integrity while producing increased stability over that of naked protoplasts. Maintenance of the cell wall in poroplasted minicells increases the osmotic resistance, mechanical resistance and storage capacity over protoplasts while permitting passage of small and medium size proteins and molecules through the porous cell wall. A poroplast is a Gram negative bacterium that has its outer membrane only removed. The production of poroplasts involves a modification of the procedure to make protoplasts to remove the outer membrane

(Birdsell et al., Production and ultrastructure of lysozyme and ethylenediaminetetraacetate-Lysozyme Spheroplasts of *Escherichia coli*, J. Bacteriology 93: 427-437, 1967; Weiss, Protoplast formation in *Escherichia coli*. J. Bacteriol. 128:668-670, 1976). Like protoplasts, measuring the total LPS that remains in the poroplast preparation may be used to monitor the removal of the outer membrane. Endotoxin kits and antibodies reactive against LPS may be used to measure LPS in solution; increasing amounts of soluble LPS indicates decreased retention of LPS by protoplasts. This assay thus makes it possible to quantify the percent removal of total outer membrane from the poroplasted minicells.

[0432] Several chemical and physical techniques have been employed to remove the outer membrane of gram negative bacteria. Chemical techniques include the use of EDTA in Tris to make cells susceptible to hydrophobic agents such as actinomycin C. Leive L. The barrier function of the gram-negative envelope. Ann NY Acad. Sci. May 10, 1974;235(0):109-29.; Voll M J, Leive L. Actinomycin resistance and actinomycin excretion in a mutant of *Escherichia coli*. J Bacteriol. 1970 May; 102(2):600-2; Lactic Acid disruption of the outer membrane as measured by the uptake of hydrophobic fluorophores; Alakomi H L, Skytta E, Saarela M, Mattila-Sandholm T, Latva-Kala K, Helander I M. Lactic acid permeabilizes gram-negative bacteria by disrupting the outer membrane. Appl Environ Microbiol. 2000 May;66(5):2001-5; and Polymyxin B disruption as measured by periplasmic constituent release (Teuber M, Cerny G. Release of the periplasmic ribonuclease I into the medium from *Escherichia coli* treated with the membrane-active polypeptide antibiotic polymyxin B. FEBS Lett. May 11, 1970;8(1):49-51). Physical techniques include the use of osmotic differentiation to facilitate the disruption of the OM. Neu H C, Heppel L A. The release of enzymes from *Escherichia coli* by osmotic shock and during the formation of spheroplasts. J. Biol. Chem. 1965 September;240(9):3685-92. See also Voll M J, Leive L. Actinomycin resistance and actinomycin excretion in a mutant of *Escherichia coli*. J Bacteriol. 1970 May;102(2):600-2; Fiil A, Branton D. Changes in the plasma membrane of *Escherichia coli* during magnesium starvation. J Bacteriol. 1969 June;98(3):1320-7; and Matsuyama S, Fujita Y, Mizushima S. SecD is involved in the release of translocated secretory proteins from the cytoplasmic membrane of *Escherichia coli*. EMBO J. 1993 January; 12(1):265-70.

[0433] III.E.3. Spheroplasts

[0434] A spheroplast is a bacterial minicell that has a disrupted cell wall and/or a disrupted OM. Unlike eubacterial minicells and poroplasts, which have a cell wall and can thus retain their shape despite changes in osmotic conditions, the absence of an intact cell wall in spheroplasts means that these minicells do not have a rigid form.

[0435] III.E.4. Protoplasts

[0436] A protoplast is a bacterium that has its outer membrane and cell wall removed. The production of protoplasts involves the use of lysozyme and high salt buffers to remove the outer membrane and cell wall (Birdsell et al., Production and ultrastructure of lysozyme and ethylenediaminetetraacetate-Lysozyme Spheroplasts of *Escherichia coli*, J. Bacteriology 93: 427-437, 1967; Weiss, Protoplast formation in *Escherichia coli*. J. Bacteriol. 128:668-670, 1976). Various commercially available lysozymes can be

used in such protocols. Measuring the total LPS that remains in the protoplast preparation is used to monitor the removal of the outer membrane. Endotoxin kits assays can be used to measure LPS in solution; increasing amounts of soluble LPS indicates decreased retention of LPS by protoplasts. This assay thus makes it possible to quantify the percent removal of total outer membrane from the minicells. Endotoxin assays are commercially available from, e.g., BioWhittaker Molecular Applications (Rockland, Me.)

**[0437]** For minicell applications that utilize bacterial-derived minicells, it may be necessary to remove the outer membrane of Gram-negative cells and/or the cell wall of any bacterial-derived minicell. For Gram-positive bacterial cells, removal of the cell wall may be easily accomplished using lysozyme. This enzyme degrades the cell wall allowing easy removal of now soluble cell wall components from the pelletable protoplasted minicells. In a more complex system, the cell wall and outer membrane of Gram-negative cells may be removed by combination treatment with EDTA and lysozyme using a step-wise approach in the presence of an osmoprotecting agent (Birdsell, et al. 1967. Production and ultrastructure of lysozyme and ethylenediaminetetraacetate-lysozyme spheroplasts of *E. coli*, J. Bacteriol. 93:427-437; Weiss, 1976. Protoplast formation in *E. coli*. J. Bacteriol. 128:668-670). By non-limiting example, this osmoprotectant may be sucrose and/or glycerol. It has been found that the concentration of the osmoprotectant sucrose, the cell wall digesting enzyme lysozyme, and chelator EDTA can be optimized to increase the quality of the protoplasts produced. Separation of either prepared Gram-negative spheroplasts prepared in either fashion from removed remaining LPS may occur through exposure of the spheroplast mixture to an anti-LPS antibody. By non-limiting example, the anti-LPS antibody may be covalently or non-covalently attached to magnetic, agarose, sepharose, sepharacyl, polyacrylamide, and/or sephadex beads. Following incubation, LPS is removed from the mixture using a magnet or slow centrifugation resulting in a protoplast-enriched supernatant.

**[0438]** Monitoring loss of LPS may occur through dot-blot analysis of protoplast mixtures or various commercially available endotoxin kit assays can be used to measure LPS in solution; increasing amounts of soluble LPS indicates decreased retention of LPS by protoplasts. This immuno assay may comprise a step of comparing the signal to a standard curve in order to quantify the percent removal of total outer membrane from the minicells. Other endotoxin assays, such as the LAL Systems from BioWhittaker, are commercially available. LPS removal has been measured by gas chromatography of fatty acid methyl esters. Alakomi H L, Skytta E, Saarela M, Mattila-Sandholm T, Latva-Kala K, Helander I M. Lactic acid permeabilizes gram-negative bacteria by disrupting the outer membrane. Appl Environ Microbiol. 2000 May;66(5):2001-5.

**[0439]** In order to reduce, preferably eliminate, in vivo antigenic potential of minicells or minicell protoplasts, minicell protoplasts may be treated to remove undesirable surface components. Minicell protoplasts so treated are referred to as "denuded minicells" a term that encompasses both spheroplasts and protoplasts. Denuding minicells or minicell protoplasts is accomplished by treatment with one or more enzymes or conditions that selectively or preferentially removes or make less antigenic externally displayed

proteins. As one non-limiting example, the protease trypsin is used to digest exposed proteins on the surface of these particles. In this example, the proteolytic activity of trypsin may be modulated or terminated by the additional of a soybean trypsin inhibitor. Non-limiting examples of other proteases that additionally or alternatively may be used include chymotrypsin, papain, elastase, proteinase K and pepsin. For some such proteases, it may be necessary to limit the extent of proteolysis by, e.g., using a suboptimal concentration of protease or by allowing the reaction to proceed for a suboptimal period of time. By the term "suboptimal," it is meant that complete digestion is not achieved under such conditions, even though the reactions could proceed to completion under other (i.e., optimal) conditions.

**[0440]** It is sometimes preferred to use molecular genetic techniques to create mutant derivatives of exogenous proteins that (1) are resistant to the proteases or other enzymes used to prepare minicells and (2) retain the desired biological activity of the receptor that is desired to be retained, i.e., the ability to bind one or more ligands of interest.

**[0441]** It is within the scope of the invention to have two or more exogenous proteins expressed within and preferentially displayed by minicells in order to achieve combined, preferable synergistic, therapeutic compositions. Similarly, two or more therapeutic minicell compositions are formulated into the same composition, or are administered during the same therapeutic minicell compositions (i.e., "cocktail" therapies). In other types of "cocktail" therapy, one or more therapeutic minicell compositions are combined or co-administered with one or more other therapeutic agents that are not minicell compositions such as, e.g., organic compounds, therapeutic proteins, gene therapy constructs, and the like.

#### III.F. Minicells from L-form Eubacteria

**[0442]** L-form bacterial strains may be used to prepare minicells and are preferred in some embodiments of the invention. L-form bacterial strains are mutant or variant strains, or eubacteria that have been subject to certain conditions, that lack an outer membrane, a cell wall, a periplasmic space and extracellular proteases. Thus, in L-form Eubacteria, the cytoplasmic membrane is the only barrier between the cytoplasm and its surrounding environment. For reviews, see Grichko, V. P., et al. 1999. The Potential of L-Form Bacteria in Biotechnology, Can. J. Chem. Engineering 77:973-977; and Gumpert J., et al. 1998 Use of cell wall-less bacteria (L-forms) for efficient expression and secretion of heterologous gene products. Curr Opin Biotechnol. 9:506-9.

**[0443]** Segregation of minicells from L-form eubacterial parent cells allows for the generation of minicells that are at least partially deficient in barriers that lie outside of the cytoplasmic membrane, thus providing direct access to components displayed on the minicell membrane. Thus, depending on the strains and methods of preparation used, minicells prepared from L-form eubacterial parent cells will be similar if not identical to various forms of poroplasts, spheroplasts and/or protoplasts. Displayed components that are accessible in L-form minicells include, but are not limited to, lipids, small molecules, proteins, sugars, nucleic acids and/or moieties that are covalently or non-covalently associated with the cytoplasmic membrane or any component thereof.

**[0444]** By way of non-limiting example, L-form Eubacteria that can be used in the methods of the invention include

species of *Escherichia*, *Streptomyces*, *Proteus*, *Bacillus*, *Clostridium*, *Pseudomonas*, *Yersinia*, *Salmonella*, *Enterococcus* and *Erwinia*. See Onoda, T., et al. 1987. Morphology, growth and reversion in a stable L-form of *Escherichia coli* K12. *J. Gen. Microbiol.* 133:527-534; Inanova, E. H., et al. 1997. Effect of *Escherichia coli* L-form cytoplasmic membranes on the interaction between macrophages and Lewis lung carcinoma cells: scanning electron microscopy. *FEMS Immunol. Med. Microbiol.* 17:27-36; Onoda, T., et al. 2000. Effects of calcium and calcium chelators on growth and morphology of *Escherichia coli* L-form NC-7. *J. Bacteriol.* 182:1419-1422; Innes, C. M., et al. 2001. Induction, growth and antibiotic production of *Streptomyces viridifaciens* L-form bacteria. *J. Appl. Microbiol.* 90:301-308; Ferguson, C. M., et al. 2000. An ELISA for the detection of *Bacillus subtilis* L-form bacteria confirms their symbiosis in strawberry. *Lett. Appl. Microbiol.* 31:390-394; Waterhouse R. N., et al. 1994. An investigation of enumeration and DNA partitioning in *Bacillus subtilis* L-form bacteria. *J. Appl. Bacteriol.* 77:497-503; Hoischen, C., et al. 2002. Novel bacterial membrane surface display system using cell wall-less L-forms of *Proteus mirabilis* and *Escherichia coli*. *Appl. Environ. Microbiol.* 68:525-531; Rippmann, J. F., et al. 1998. Prokaryotic expression of single-chain variable-fragment (scFv) antibodies: secretion in L-form cells of *Proteus mirabilis* leads to active product and overcomes the limitations of periplasmic expression in *Escherichia coli*. *Appl. Environ. Microbiol.* 64:4862-4869; Mahony, D. E., et al. 1988. Transformation of *Clostridium perfringens* L forms with shuttle plasmid DNA. *Appl. Environ. Microbiol.* 54:264-267; Kurona, M., et al. 1983. Intergenous cell fusions between L-form cells of *Pseudomonas aeruginosa* and *Escherichia coli*. *Biken. J.* 26:103-111; Ivanova, E., et al. 2000. Studies of the interactions of immunostimulated macrophages and *Yersinia enterocolitica* 0:8. *Can. J. Microbiol.* 46:218-228; Allan, E. J., et al. 1993. Growth and physiological characteristics of *Bacillus subtilis* L-forms. *J. Appl. Bacteriol.* 74:588-594; Allan, E. J. 1991. Induction and cultivation of a stable L-form of *Bacillus subtilis*. *J. Appl. Bacteriol.* 70:339-343; Nishikawa, F., et al. 1994. Protective capacity of L-form *Salmonella typhimurium* against murine typhoid in C3H/HeJ mice. *Microbiol. Immunol.* 38:129-137; Kita, E., et al. 1993. Isolation of a cytotoxin from L-form *Salmonella typhimurium*. *FEMS Microbiol. Lett.* 109:179-184; Jass, J., et al. Growth and adhesion of *Enterococcus faecium* L-forms. *FEMS Microbiol. Lett.* 115:157-162; and U.S. Pat. No. 6,376,245.

#### [0445] IV. Assaying Minicells

##### [0446] IV.A. Efficiency of Minicell Production

[0447] The level of minicell production will vary and may be evaluated using methods described herein. Relatively high levels of minicell production are generally preferred. Conditions in which about 40% of cells are achromosomal have been reported (see, e.g., Hassan et al., Suppression of initiation defects of chromosome replication in *Bacillus subtilis* dnaA and oriC-deleted mutants by integration of a plasmid replicon into the chromosomes, *J. Bacteriol.* 179:2494-502, 1997). Procedures for identifying strains that give high yields of minicells are known in the art; see, e.g., Clark-Curtiss and Curtiss III, Analysis of Recombinant DNA Using *Escherichia coli* Minicells, *Meth. Enzol.* 101:347-362, 1983.

[0448] Minicell production can be assessed by microscopic examination of late log-phase cultures. The ratio of minicells to normal cells and the frequency of cells actively producing minicells are parameters that increase with increasing minicell production.

##### [0449] IV.B. Detecting Protein Synthesis in Minicells

[0450] Methods for detecting and assaying protein production are known in the art. See, e.g., Clark-Curtiss and Curtiss III, *Meth. Enzol.* 101:347-362, 1983. As an exemplary procedure, transformed *E. coli* minicell-producing cells are grown in LB broth with the appropriate antibiotic overnight. The following day the overnight cultures are diluted 1:50 in fresh media, and grown at 37° C. to mid-log phase. If it is desired to eliminate whole cells, an antibiotic that kills growing (whole) cells but not quiescent cells (minicells) may be used. For example, in the case of cells that are not ampicillin resistant, ampicillin (100 mg per ml is added), and incubation is allowed to continue for about 2 more hrs. Cultures are then centrifuged twice at low speed to pellet most of the large cells. Minicells are pelleted by spinning 10 min at 10,000 rpm, and are then resuspended in M63 minimal media supplemented with 0.5% casamino acids, and 0.5 mM cAMP, or M9 minimal medium supplemented with 1 mM MgSO<sub>4</sub>, 0.1 mM CaCl<sub>2</sub>, 0.05% NaCl, 0.2% glucose, and 1 ng per ml thiamine. Labeled (<sup>35</sup>S) methionine is added to the minicells for about 15 to about 90 minutes, and minicells are immediately collected afterwards by centrifugation for 10 min at 4° C. and 14,000 rpm. Cells are resuspended in 50 to 100 μg Laemmli-buffer, and disrupted by boiling and vortexing (2 min for each step). Incorporation of <sup>35</sup>S-methionine was determined by measuring the amount of radioactivity contained in 1 μl of the lysate after precipitation of proteins with trichloroacetic acid (TCA). Minicell lysates (50,000 to 100,000 cpm per lane) are subjected to PAGE on, e.g., 10% polyacrylamide gels in which proteins of known size are also run as molecular weight standards. Gels are fixed and images there of are generated by, e.g., autoradiography or any other suitable detection systems.

##### [0451] IV.C. Evaluating the Therapeutic Potential of Minicells

[0452] Various methods are used at various stages of development of a therapeutic minicell composition to estimate their therapeutic potential. As a non-limiting example, the therapeutic potential of minicells displaying a receptor is examined as follows.

##### [0453] IV.C.1. Receptors

[0454] The specificity of, rate of association of, rate of dissociation of, and/or stability of complexes resulting from, receptor binding to its ligand can be measured in vitro using methods known in the art.

[0455] In the case of a sphingolipid binding receptor, such as an S1P receptor, the ligand (S1P) is detectably labeled so that the specificity of, rate of formation of, and degree of stability of complexes resulting from the ligand-receptor binding can be examined by measuring the degree and rate at which the labeled ligand is removed from solution due to its binding to minicells displaying the receptor. In order to avoid extraneous factors from influencing these experiments, they are carried out in buffered solutions that are as free of contaminating substances as possible. However, as is understood in the art, stabilizing agents such as BSA (bovine

serum albumin) or protease inhibitors may be desirably included in these experiments. In a preferred environment, a sphingolipid binding receptor is the rat EDG-1, rat EDG-3, rat SCaMPER and human SCaMPER, the sequences of which are set forth herein.

[0456] Minicell compositions that bind sphingolipids with the desired specificity are identified from the preceding experiments. Typically, studies of ligand-receptor binding then proceed to studies in which the binding capacity of promising minicell compositions is tested under in vitro conditions that are increasingly more representative of in vivo conditions. For example, binding experiments are carried out in the presence of sera or whole blood in order to determine the therapeutic potential of minicell compositions in the presence of compounds that are present within the circulatory system of an animal.

#### [0457] IV. C.2. Molecular Sponge

[0458] Minicell compositions can also be tested for their ability to bind and/or internalize toxic compounds. The therapeutic potential of such capacity is evaluated using experiments in which detectably labeled derivatives of a toxic compound are present in the bloodstream of an anesthetized animal, which may be a human. The blood of the animal is shunted out of the body and past a device that incorporates a minicell composition before being returned to the body. The device is constructed so that the blood contacts a semipermeable membrane that is in contact with the minicell composition. By "semipermeable" it is meant that certain agents can be freely exchanged across the membrane, whereas others are retained on one side of the membrane or the other. For example, the toxic compound of interest is able to cross the semipermeable membrane, whereas minicells and blood cells are separately retained in their respective compartments. Detectably labeled derivatives of the toxic compound are present in the bloodstream of the animal. The capacity of the minicells to take up the toxic compound corresponds with a reduction of the levels of detectably labeled material in the blood and an increase in detectably labeled material in the minicell composition.

[0459] The above types of minicell-comprising compositions, devices, and procedures may be incorporated into ex vivo modalities such as ex vivo gene therapy and dialysis machines. An "ex vivo modality" is one in which a biological sample, such as a blood sample, is temporarily removed from an animal, altered through in vitro manipulation, and then returned to the body. In "ex vivo gene therapy," cells in the sample from the animal are transformed with DNA in vitro and then returned to the body. A "dialysis machine" is a device in which a fluid such as blood of an animal is temporarily removed therefrom and processed through one or more physical, chemical, biochemical, binding or other processes designed to remove undesirable substances including but are not limited to toxins, venoms, overexpressed or overactive endogenous agents, and pathogens or molecules derived therefrom.

[0460] Intraminicellular co-expression of a second molecule that is displayed on the surface of minicells, and which is a ligand for a binding moiety that is immobilized, can optionally be used in order to remove minicells from the sample before it is returned to the body. In the latter aspect, minicells that bind undesirable substances are preferably removed with the undesirable compound remaining bound

to the minicells. Minicells that have been used for ex vivo gene therapy, but which have failed to deliver a nucleic acid to any cells in the sample, can be removed in a similar manner.

#### [0461] IV.C.3. Minicell-Solubilized Receptors

[0462] It is known in the art to use recombinant DNA technology to prepare soluble (hydrophilic) receptor fragments from receptors that bind a bioactive ligand. Unlike the native, membrane-bound receptor, which is relatively insoluble in water (hydrophobic), soluble receptor fragments can be formulated for therapeutic delivery using techniques that are known to have been used to formulate soluble agents.

[0463] Typically, soluble receptor fragments are used to competitively inhibit the binding of the receptor to its ligand. That is, the soluble receptor fragments bind the ligand at the expense of the membrane-bound receptor. Because less of the ligand is bound to its receptor, the cellular response to the ligand is attenuated. Common cellular responses that are desirably attenuated include but are not limited to the uptake of an undesirable agent (e.g., a toxin, a pathogen, etc.) and activation of a signaling pathway having undesirable consequences (e.g., inflammation, apoptosis, unregulated growth, etc.).

[0464] Preparing a soluble fragment derived from a receptor is not trivial. Typically, the three dimensional structure of the receptor is not known, and must be predicted based on homology with other receptors or by using software that predicts the tertiary structure of a polypeptide based on its amino acid sequence. Using the hypothetical structure of the receptor, a series of polypeptides are prepared that comprise amino acid sequences from the receptor but lack regions thereof that are thought to be membrane-anchoring or transmembrane domain(s) of the receptor. Some of the polypeptides prepared this way may be soluble, some may retain the binding activity of the receptor, and a few may have both characteristics. Members of the latter class of polypeptides are soluble receptor fragments, some of which may be amenable to development as a therapeutic or diagnostic agent.

[0465] For any given receptor, there is always the possibility that none of the soluble fragments derived from the receptor will specifically bind its ligand with sufficient affinity as to be therapeutically effective. Thus, in some instances, it may not be possible to prepare a receptor fragment that is both soluble and sufficiently biologically active.

[0466] The minicells of the invention provide a "universal carrier" for receptors that allows the hydrophobic receptors to be solubilized in the sense that, although they remain associated with a membrane, the minicell is a small, soluble particle. That is, as an alternative to preparing a set of polypeptides to see which, if any of them, are water soluble receptor fragments, one may, using the teachings of the disclosure, prepare soluble minicells that display the receptor.

#### [0467] IV.C.4. Reducing Toxicity

[0468] For in vivo use of minicells for the purposes of eliciting an immune response or for therapeutic and diagnostic applications involving delivery of minicells to a

human or to an animal, it may be useful to minimize minicell toxicity by using endotoxin-deficient mutants of parent cells. Without being limited to the following example, lipopolysaccharide (LPS) deficient *E. coli* strains could be conjugated with minicell producing cells to make parent cells lacking the endotoxin. LPS synthesis in *E. coli* includes the lipid A biosynthetic pathway. Four of the genes in this pathway have now been identified and sequenced, and three of them are located in a complex operon which also contains genes involved in DNA and phospholipid synthesis. The *rfa* gene cluster, which contains many of the genes for LPS core synthesis, includes at least 17 genes. The *rfb* gene cluster encodes protein involved in O-antigen synthesis, and *rfb* genes have been sequenced from a number of serotypes and exhibit the genetic polymorphism anticipated on the basis of the chemical complexity of the O antigens (Schnaitman and Klena. 1993. Genetics of lipopolysaccharide biosynthesis in enteric bacteria. *Microbiol. Rev.* 57:655-82). When present alone or in combination the *rfb* and *oms* mutations cause alterations in the eubacterial membrane that make it more sensitive to lysozyme and other agents used to process minicells. Similarly, the *rfa* (Chen, L., and W. G. Coleman Jr. 1993. Cloning and characterization of the *Escherichia coli* K-12 *rfa-2* (*rfaC*) gene, a gene required for lipopolysaccharide inner core synthesis. *J. Bacteriol.* 175:2534-2540), *IpcA* (Brooke, J. S., and M. A. Valvano. 1996. Biosynthesis of inner core lipopolysaccharide in enteric bacteria identification and characterization of a conserved phosphoheptose isomerase. *J. Biol. Chem.* 271:3608-3614), and *IpcB* (Kadrman, J. L., et al. 1998. Cloning and overexpression of glycosyltransferases that generate the lipopolysaccharide core of *Rhizobium leguminosarum*. *J. Biol. Chem.* 273:26432-26440) mutations, when present alone or in combination, cause alterations in lipopolysaccharides in the outer membrane causing cells to be more sensitive to lysozyme and agents used to process minicells. In addition, such mutations can be used to reduce the potential antigenicity and/or toxicity of minicells.

[0469] Minicell-producing cells may comprise mutations that augment preparative steps. For example, lipopolysaccharide (LPS) synthesis in *E. coli* includes the lipid A biosynthetic pathway. Four of the genes in this pathway have now been identified and sequenced, and three of them are located in a complex operon that also contains genes involved in DNA and phospholipid synthesis. The *rfa* gene cluster, which contains many of the genes for LPS core synthesis, includes at least 17 genes. The *rfb* gene cluster encodes protein involved in O-antigen synthesis, and *rfb* genes have been sequenced from a number of serotypes and exhibit the genetic polymorphism anticipated on the basis of the chemical complexity of the O antigens. See Schnaitman and Klena, Genetics of lipopolysaccharide biosynthesis in enteric bacteria, *Microbiol. Rev.* 57:655-82, 1993. When present, alone, or in combination, the *rfb* and *oms* mutations cause alterations in the eubacterial membrane that make it more sensitive to lysozyme and other agents used to process minicells. Similarly, the *rfa* (Chen, L., and W. G. Coleman Jr. 1993. Cloning and characterization of the *Escherichia coli* K-12 *rfa-2* (*rfaC*) gene, a gene required for lipopolysaccharide inner core synthesis. *J. Bacteriol.* 175:2534-2540), *IpcA* (Brooke, J. S., and M. A. Valvano. 1996. Biosynthesis of inner core lipopolysaccharide in enteric bacteria identification and characterization of a conserved phosphoheptose isomerase. *J. Biol. Chem.* 271:3608-3614), and *IpcB*

(Kadrman, J. L., et al. 1998. Cloning and overexpression of glycosyltransferases that generate the lipopolysaccharide core of *Rhizobium leguminosarum*. *J. Biol. Chem.* 273:26432-26440) mutations, when present alone or in combination, cause alterations in lipopolysaccharides in the outer membrane causing cells to be more sensitive to lysozyme and agents used to process minicells. In addition, such mutations can be used to reduce the potential antigenicity and/or toxicity of minicells.

#### [0470] V. Genetic Expression in Minicells

[0471] Various minicells of the invention use recombinant DNA expression systems to produce a non-eubacterial protein, which may be a membrane protein that is preferably "displayed" on the surface of minicells, a membrane protein that projects portions not associated with a membrane towards the interior of a minicell, or a soluble protein present in the exterior of the minicells. By "displayed" it is meant that a protein is present on the surface of a cell (or minicell) and is thus in contact with the external environment of the cell. Non-limiting examples of displayed exogenous proteins of the invention include mammalian receptors and fusion proteins comprising one or more transmembrane domains. In other aspects of the invention, minicells use expression elements to produce bioactive nucleic acids from templates therefor.

#### [0472] V.A. Expression Systems

[0473] In vivo and in vitro protein expression systems provide a variety of techniques that allow scientists to transcribe and translate amino acid polypeptides proteins from recombinant DNA templates (Kaufman, Overview of vector design for mammalian gene expression. *Mol. Biotechnol.* 2001. 16: 151-160; and Kozak, Initiation of translation in prokaryotes and eukaryotes. *Gene*, 1999. 234: 187-208).

[0474] Although minicells are virtually depleted of chromosomal DNA (Tudor et al., Presence of nuclear bodies in some minicells of *Escherichia coli*. *J. Bacteriol.* 1969. 98: 298-299), it has been reported that minicells have all the elements required to express nucleotide sequences that are present in episomal expression elements therein (Levy, Very stable prokaryote messenger RNA in chromosomeless *Escherichia coli* minicells. *Proc Natl Acad Sci USA*, 1975. 72: 2900-2904; Hollenberg et al., Synthesis of high molecular weight polypeptides in *Escherichia coli* minicells directed by cloned *Saccharomyces cerevisiae* 2-micron DNA. *Gene*, 1976. 1: 33-47; Crooks et al., Transcription of plasmid DNA in *Escherichia coli* minicells. *Plasmid*, 1983. 10: 66-72; Clark-Curtiss, Analysis of recombinant DNA using *Escherichia coli* minicells. *Methods Enzymol.* 1983. 101: 347-362).

[0475] Preferred expression vectors and constructs according to the invention are episomal genetic elements. By "episomal" it is meant that the expression construct is not always linked to a cell's chromosome but may instead be retained or maintained in host cells as a distinct molecule entity. Minicells can retain, maintain and express episomal expression constructs such as, e.g., plasmids, bacteriophage, viruses and the like (Crooks et al., *Plasmid* 10:66-72, 1983; Clark-Curtiss, *Methods Enzymology* 101:347-62, 1983; Witkiewicz et al., *Acta. Microbiol. Pol.* A 7:21-24, 1975; Ponta et al., *Nature* 269:440-2, 1977). By "retained" it is

meant that the episomal expression construct is at least temporarily present and expressed in a host parent cell and/or minicell; by "maintained" it is meant that the episomal expression construct is capable of autonomous replication within a host parent cell and/or minicell. In the context of episomal elements, the term "contained" encompasses both "retained" and "maintained." A preferred type of an episomal element according to the invention is one that is always an extrachromosomal element, or which is part of a chromosome but becomes an extrachromosomal element before or during minicell production.

**[0476]** The fact that minicells do not contain chromosomal DNA but do contain episomal expression elements, such as plasmids, that can be used as templates for RNA synthesis means that the only proteins that are actively produced in minicells are those that are encoded by the expression elements that they contain. Minicell-producing *E. coli* cells can be made competent and transformed with expression elements that direct the expression of proteins encoded by the expression elements. An expression element segregates into minicells as they are produced. In isolated minicells that contain expression elements, there is a single DNA template RNA for transcription. Therefore, the only nucleic acids and proteins that are actively produced (expressed) by minicells are those that are encoded by sequences on the expression vector. In the context of the invention, sequences that encode amino acid sequences are designated "open reading frames" or "ORFs." One feature of minicell expression systems of interest as regards the present invention is that endogenous (i.e., chromosomally located) genes are not present and are thus not expressed, whereas genes present on the episomal element are expressed (preferably over-expressed)-in the minicells. As a result, the amount of endogenous proteins, including membrane proteins, decreases as the minicells continue to express genes located on episomal expression constructs.

**[0477]** The minicell system can reduce or eliminate undesirable features associated with the transcription and translation of endogenous proteins from the *E. coli* chromosome. For example, expression of proteins in minicell systems results in low background signal ("noise") when radiolabeled proteins produced using recombinant DNA technology (Jannatipour et al., Translocation of Vibrio Harveyi N,N'-Diacetylchitinase to the outer membrane of *Escherichia coli*. J. Bacteriol, 1987. 169: 3785-3791). A high background signal can make it difficult to detect a protein of interest. In whole cell *E. coli* systems, endogenous proteins (encoded by the bacterial chromosome) are labeled as well as the protein(s) encoded by the expression element; whereas, in minicell systems, only the proteins encoded by the expression element in the minicells are labeled.

**[0478]** There are a variety of proteins, both eubacterial and eukaryotic, that have been expressed from plasmid DNA in minicells (Clark-Curtiss, Methods Enzymol, 101:347-362, 1983). Some examples of proteins and nucleic acids that have been expressed in minicells include the Kdp-ATPase of *E. coli* (Altendorf et al., Structure and function of the Kdp-ATPase of *Escherichia coli*. Acta Physiol Scand, 643: 137-146, 1998); penicillin binding proteins alpha and gamma (Davies et al., Prediction of signal sequence-dependent protein translocation in bacteria: Assessment of the *Escherichia coli* minicell system. Biochem Biophys Res Commun, 150: 371-375, 1988); cell surface antigens of

*Polyomaonas gingivalis* (Rigg et al., The molecular cloning, nucleotide sequence and expression of an antigenic determinant from *Porphyromonas gingivalis*. Arch Oral Biol, 45:41-52, 2000); trkG integral membrane protein of *E. coli* (Schlosser et al., Subcloning, Nucleotide sequence, and expression of trkG, a gene that encodes an integral membrane protein involved in potassium uptake via the Trk system of *Escherichia coli*. J. Bacteriol, 173:3170-3176, 1991); the 34 kDa antigen of *Treponema pallidum* (Swancutt et al., Molecular characterization of the pathogen-specific, 34-kilodalton membrane immunogen of *Treponema pallidum*. Infect Immun, 57:3314-23, 1989); late proteins of bacteriophage MB78 (Colla et al., IUBMB Life 48:493-497, 1999); uncharacterized DNA from *Xenopus laevis* (Cohen and Boyer, U.S. Pat. No. 4,237,224, which issued Dec. 2, 1980); the onc gene v-fos (MacConnell and Verman, Expression of FBJ-MSV oncogene (fos) product in bacteria, 131(2) Virology 367 1983); interferon (Edge et al., Chemical synthesis of a human interferon-alpha 2 gene and its expression in *Escherichia coli*, Nucleic Acids Res. 11:6419, 1983); bovine growth hormone (Rosner et al., Expression of a cloned bovine growth hormone gene in *Escherichia coli* minicells, Can. J. Biochem. 60:521-4, 1982); gastrointestinal hormone (Suzuki et al., Production in *Escherichia coli* of biologically active secretin, a gastrointestinal hormone, Proc. Natl. Acad. Sci. USA 79:2475, 1982); and archaeobacterial proteins (Lienard and Gottschalk, Cloning, sequencing and expression of the genes encoding the sodium translocating N-methyltetrahydromethanopterin: coenzyme M methyltransferase of the methylotrophic archaeon *Methanosarcina mazei* GoI, 425 FEBS Letters 204, 1998; and Lemker et al., Overproduction of a functional Al ATPase from the archaeon *Methanosarcina mazei* GI in *Escherichia coli*, European Journal of Biochemistry 268:3744, 2001).

**[0479]** V.B. Modulating Genetic Expression in Minicells

**[0480]** Gene expression in minicells, and/or in minicell-producing (parent) cells, involves the coordinated activity of a variety of expression factors, regulatory elements and expression sequences. Any of these may be modified to alter the extent, timing or regulation of expression of a gene of interest in minicells and/or their parent cells. Often, the goal of the manipulations is to increase the efficiency of protein production in minicells. However, increased expression may, in some instances, desirably include increased or "tight" negative regulation. This may reduce or eliminate selective pressure created by toxic gene products, and allow for functional expression in a controlled fashion by removing the negative regulation and/or inducing expression of the gene product at a preselected time. By way of non-limiting example, these techniques may include modification or deletion of endogenous gene(s) from which their respective gene product decreases the induction and expression efficiency of a desired protein in the parent cell prior to minicell formation and/or the segregated minicell. By way of non-limiting example, these protein components may be the enzymes that degrade chemical inducers of expression, proteins that have a dominant negative affect upon a positive regulatory elements, proteins that have proteolytic activity against the protein to be expressed, proteins that have a negative affect against a chaperone that is required for proper activity of the expressed protein, and/or this protein may have a positive effect upon a protein that either degrades or prevents the proper function of the expressed protein. These gene products that require deletion or modi-



fication for optimal protein expression and/or function may also be antisense nucleic acids that have a negative affect upon gene expression.

[0481] VI. Fusion (Chimeric) Proteins

[0482] In certain aspects of the invention, a fusion protein is expressed and displayed by minicells. One class of fusion proteins of particular interest are those that are displayed on the surface of minicells, e.g., fusion proteins comprising one or more transmembrane domains. Types of displayed fusion proteins of particular interest are, by way of non-limiting example, those that have an extracellular domain that is a binding moiety or an enzymatic moiety. By way of non-limiting example, the fusion protein ToxR-PhoA has been expressed in and displayed on the surface of minicells. The ToxR-PhoA fusion protein comprises a polypeptide corresponding to the normally soluble enzyme, alkaline phosphatase, anchored to the minicell membrane by the single transmembrane domain of ToxR (see the Examples). The fusion protein retains the activity of the enzyme in the context of the minicell membrane in which it is bound. Nearly all of the fusion protein is oriented so that the enzyme's catalytic domain is displayed on the outer surface of the minicell.

[0483] VI.A. Generation of Fusion Proteins

[0484] Polypeptides, which are polymers of amino acids, are encoded by another class of molecules, known as nucleic acids, which are polymers of structural units known as nucleotides. In particular, proteins are encoded by nucleic acids known as DNA and RNA (deoxyribonucleic acid and ribonucleic acid, respectively).

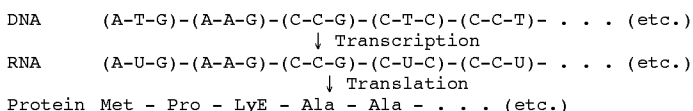
[0485] The nucleotide sequence of a nucleic acid contains the "blueprints" for a protein. Nucleic acids are polymers of nucleotides, four types of which are present in a given nucleic acid. The nucleotides in DNA are adenine, cytosine and guanine and thymine, (represented by A, C, G, and T respectively); in RNA, thymine (T) is replaced by uracil (U). The structures of nucleic acids are represented by the sequence of its nucleotides arranged in a 5' ("5 prime") to 3' ("3 prime") direction, e.g.,

[0486] 5-A-T-G-C-C-T-A-A-A-G-C-C-G-C-T-C-C-C-T-C-A-3'

[0487] In biological systems, proteins are typically produced in the following manner. A DNA molecule that has a nucleotide sequence that encodes the amino acid sequence of a protein is used as a template to guide the production of a messenger RNA (mRNA) that also encodes the protein; this process is known as transcription. In a subsequent process called translation, the mRNA is "read" and directs the synthesis of a protein having a particular amino acid sequence.

[0488] Each amino acid in a protein is encoded by a series of three contiguous nucleotides, each of which is known as a codon. In the "genetic code," some amino acids are encoded by several codons, each codon having a different sequence; whereas other amino acids are encoded by only one codon sequence. An entire protein (i.e., a complete amino acid sequence) is encoded by a nucleic acid sequence called a reading frame. A reading frame is a continuous nucleotide sequence that encodes the amino acid sequence of a protein; the boundaries of a reading frame are defined by its initiation (start) and termination (stop) codons.

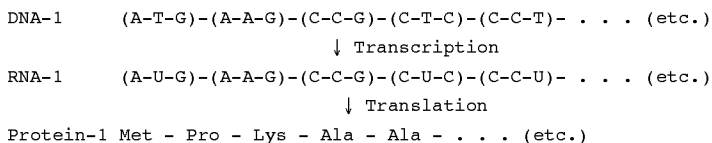
[0489] The process by which a protein is produced from a nucleic acid can be diagrammed as follows:



[0490] A chimeric reading frame encoding a fusion protein is prepared as follows. A "chimeric reading frame" is a genetically engineered reading frame that results from the fusion of two or more normally distinct reading frames, or fragments thereof, each of which normally encodes a separate polypeptide. Using recombinant DNA techniques, a first reading frame that encodes a first amino acid sequence is linked to a second reading frame that encodes a second amino acid sequence in order to generate a chimeric reading frame. Chimeric reading-frames may also include nucleotide sequences that encode optional fusion protein elements (see below).

[0491] A hypothetical example of a chimeric reading frame created from two normally separate reading frames is depicted in the following flowchart.

[0492] First Open Reading Frame and "Protein-1":



**[0493]** Second Open Reading Frame and "Protein-2":

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DNA-2      (T-G-G)-(G-T-T)-(A-C-T)-(C-A-C)-(T-C-A)- . . . (etc.)
              ↓ Transcription
RNA-2      (U-G-G)-(G-U-U)-(A-C-U)-(C-A-C)-(U-C-A)- . . . (etc.)
              ↓ Translation
Protein-2  Trp - Val - Thr - His - Ser - . . . (etc.)

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**[0494]** Chimeric Reading Frame that encodes a Fusion Protein having sequences from Protein-1 and Protein-2:

example of the former class of detectable polypeptide is green fluorescent protein (GFP). Examples of the latter class

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DNA-Chimera (A-T-G)-(A-A-G)-(C-C-G)-(C-A-C)-(T-C-A)-(etc.)
              ↓ Transcription
RNA-Chimera (A-U-G)-(A-A-G)-(C-C-G)-(C-A-C)-(U-C-A)-(etc.)
              ↓ Translation
Fusion Protein Met - Pro - Lys - His - Ser - (etc.)

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**[0495]** In order for a chimeric reading frame to be functional, each normally distinct reading frame therein must be fused to all of the other normally distinct reading frames in a manner such that all of the reading frames are in frame with each other. By "in frame with each other" it is meant that, in a chimeric reading frame, a first nucleic acid having a first reading frame is covalently linked to a second nucleic acid having a second reading frame in such a manner that the two reading frames are "read" (translated) in register with each other. As a result, the chimeric reading frame encodes one extended amino acid sequence that includes the amino acid sequences encoded by each of the normally separate reading frames. A fusion protein is thus encoded by a chimeric reading frame.

**[0496]** The fusion proteins of the invention are used to display polypeptides on minicells. The fusion proteins comprise (1) at least one polypeptide that is desired to be displayed by minicells (a "displayed polypeptide") and (2) at least one membrane polypeptide, e.g., a transmembrane or a membrane anchoring domain. For various aspects of the invention, optional fusion protein elements, as defined herein, may also be included if required or desired.

**[0497]** VI.B. Optional Fusion Protein Elements

**[0498]** The fusion proteins of the invention may optionally comprise one or more non-biologically active amino acid sequences, i.e., optional fusion protein elements. Such elements include, but are not limited to, the following optional fusion protein elements. It is understood that a chimeric reading frame will include nucleotide sequences that encode such optional fusion protein elements, and that these nucleotide sequences will be positioned so as to be in frame with the reading frame encoding the fusion protein. Optional fusion protein elements may be inserted between the displayed polypeptide and the membrane polypeptide, upstream or downstream (amino proximal and carboxy proximal, respectively) of these and other elements, or within the displayed polypeptide and the membrane polypeptide. A person skilled in the art will be able to determine which optional element(s) should be included in a fusion protein of the invention, and in what order, based on the desired method of production or intended use of the fusion protein.

**[0499]** Detectable polypeptides are optional fusion protein elements that either generate a detectable signal or are specifically recognized by a detectably labeled agent. An

include epitopes such as a "His tag" (6 contiguous His residues, a.k.a. 6x His), the "FLAG tag" and the c-myc epitope. These and other epitopes can be detected using labeled antibodies that are specific for the epitope. Several such antibodies are commercially available.

**[0500]** Attachment (support-binding) elements are optionally included in fusion proteins and can be used to attach minicells displaying a fusion protein to a preselected surface or support. Examples of such elements include a "His tag," which binds to surfaces that have been coated with nickel; streptavidin or avidin, which bind to surfaces that have been coated with biotin or "biotinylated" (see U.S. Pat. No. 4,839,293 and Airene et al., *Protein Expr. Purif.* 17:139-145, 1999); and glutathione-s-transferase (GST), which binds to surfaces coated with glutathione (Kaplan et al., *Protein Sci.* 6:399-406, 1997; U.S. Pat. No. 5,654,176). Polypeptides that bind to lead ions have also been described (U.S. Pat. No. 6,111,079).

**[0501]** Spacers (a.k.a. linkers) are amino acid sequences that are optionally included in a fusion protein in between other portions of a fusion protein (e.g., between the membrane polypeptide and the displayed polypeptide, or between an optional fusion protein element and the remainder of the fusion protein). Spacers can be included for a variety of reasons. For example, a spacer can provide some physical separation between two parts of a protein that might otherwise interfere with each other via, e.g., steric hindrance. The ability to manipulate the distance between the membrane polypeptide and the displayed polypeptide allows one to extend the displayed polypeptide to various distances from the surface of minicells.

**[0502]** VI.C. Interactions with Receptient Cells

**[0503]** Many Gram-negative pathogens use a type III secretion machine to translocate protein toxins across the bacterial cell envelope (for a review, see Cheng L W, Schneewind O. Type III machines of Gram-negative bacteria: delivering the goods. *Trends Microbiol* 2000 May;8(5):214-20). For example, pathogenic *Yersinia* spp. export over a dozen Yop proteins via a type III mechanism, which recognizes secretion substrates by signals encoded in yop mRNA or chaperones bound to unfolded Yop proteins. A 70-kb virulence plasmid found in pathogenic *Yersinia* spp. to survive and multiply in the lymphoid tissues of the host. The virulence plasmid encodes the Yop virulon, an integrated system allowing extracellular bacteria to inject bac-

terial proteins into cells. The Yop virulon comprises a variety of Yop proteins and a dedicated type III secretion apparatus, called Ysc (for a review, see Cornelis G R, Boland A, Boyd A P, Geuijen C, Iriarte M, Neyt C, Sory M P, Stainier I. The virulence plasmid of *Yersinia*, an antihost genome. *Microbiol Mol Biol Rev* 1998 62(4):1315-52).

#### [0504] VII. Minicell Display

[0505] Included in the design of the invention is the use of minicells to express and display soluble or membrane-bound protein libraries to identify a soluble or membrane-bound protein that binds a known ligand or to identify proteins (e.g. orphan receptors) for which the known ligand or substrate is not known but for which a reporter could be engineered into the minicell that would signal the presence of the encoded protein. In the preferred embodiment of the invention, this 'minicell display' technique is analogous to phage display for the purpose of identifying genes that encode receptor-like or antibody-like proteins against known ligand. This approach will allow identification of an unknown receptor protein for which a known ligand has affinity. These known ligands may have been identified as having a pharmacological, biological, or other effect without knowledge of the site of effect. In these cases the knowledge of receptor will allow basic research to understand the molecular and/or physiological response and permit directed modification of the ligand for better pharmacological or biological response or modification of the receptor for employment in ligand-binding applications. In another non-limiting embodiment of the invention, the ligand need not be known but some general characteristic of the protein would be.

[0506] For purposes of this application, soluble or membrane-bound protein libraries may be constructed by random cloning of DNA fragments or directed cloning using reverse transcriptase polymerase chain reaction (RT-PCR). In either method, DNA fragments may be placed under the regulation of any regulatory element listed in section II.B. on any plasmid or chromosomal construct. In the case of soluble protein receptors, they will be fused to form a chimeric protein with a known transmembrane domain (TMD), e.g. the TMD from the *toxR* gene product. Upon induction of the soluble or membrane-bound protein library, minicells, minicell protoplasts, or minicell poroplasts (as the experiment requires) will be mixed with the known ligand. Without being limited to the following example, screening could be accomplished by first labeling the known ligand with a molecular fluorophore, e.g. TAMRA, FTC, or in some cases a fluorescent protein, e.g. GFP. A positive interaction between the minicells displaying the receptor for the labeled ligand will be identified and separated from the library population by fluorescent-activated cell sorting (FACS). Isolated, positive receptor-ligand interactions will be identified by PCR amplification, subcloned into a clean background, and sequenced using plasmid-specific oligonucleotides. Subcloned proteins will be re-screened for interaction with the labeled ligand, and their binding patterns characterized.

[0507] Positive interacting receptor proteins may be employed in mutagenesis or other directed evolutionary process to improve or decrease the binding affinity to the ligand. In another application, the receptor-ligand pair may be further employed in a screening process to identify new compounds that may interfere with the interaction. Thus,

using a known substance to identify the receptor and the identified receptor-ligand pair to identify other interfering compounds. Chimeric-soluble or membrane-bound protein libraries may be screened versus a protein-array chip that presents a variety of known protein compounds or peptide variations. In this application, the minicell, minicell protoplast, or minicell poroplast will also contain a label, signaling component, and/or antigen recognizable by an antibody for identification of a positive interaction on the protein chip array. Other approaches for identification may include packaged fluorescent molecules or proteins that are constitutively produced, induced by the positive interaction with the ligand, or regulated by a regulatory element described in section II.B.

[0508] In a preferred embodiment of the invention, cDNA libraries could be constructed from isolated B-cells, activated B-cell or T-cells for the purpose of identifying receptors or antibodies that are encoded by these cells of the immune system. In a non-limiting example, a small molecule could be used to immunize an experimental animal (e.g., rat, mouse, rabbit), the spleen could be removed, or blood could be drawn and used as a source of mRNA. Reverse transcription reactions could then be used to construct a cDNA library that would eventually be transformed into the minicell parent bacteria, as described above. The minicells would then be isolated, induced and subjected to FACS analysis with subsequent amplification and sequencing of the cDNA fragment of interest (see above). The PCR-amplified plasmid-containing cDNA fragment encoding the "receptor" or "antibody" of interest would be ready for transformation and expression in the minicell context for diagnostic, therapeutic research or screening applications of the invention.

[0509] In a related, non-limiting embodiment of the invention, minicells expressing a particular antigen (e.g., protein, carbohydrate, small molecule, lipid) on their surfaces (described elsewhere in this application) are used to generate an immunogenic response. The advantages of presenting an antigen on the surfaces of minicells are that the minicells themselves may be an adjuvant that stimulates the immune response, particularly if administered subcutaneously (SC) or intramuscularly (IM). Moreover, the minicells are not readily eliminated by the renal system and are present in the circulatory system of an immunized animal for a longer time. In addition, small molecules could be tethered to the minicell in a way that presents the desired moiety of the molecule. Animals are presented with minicell-based immunogens, and the antibodies produced in the animals are prepared and used in therapeutic, diagnostic, research and screening applications. Although this aspect of the invention may, be used to make antibodies to any molecule displayed on their surface, the extracellular domains of membrane proteins are of particular interest.

[0510] Minicell display could be used to identify orphan receptors or other proteins for which a ligand or substrate is not known. As a non-limiting example, orphan G protein coupled receptors (GPCRs) or novel RNA and DNA polymerases could be identified from organisms living in extreme environments. A cDNA library could be constructed from an organism and expressed in minicells that co-express a reporter system that indicates the presence of the novel protein. In a non-limiting example of GPCRs, the

minicells used for minicell display are engineered to express a G-protein in a manner that would signal an interaction with the orphan GPCR.

**[0511]** VIII. Aptamers

**[0512]** Traditionally, techniques for detecting and purifying target molecules have used polypeptides, such as antibodies, that specifically bind such targets. While nucleic acids have long been known to specifically bind other nucleic acids (e.g., ones having complementary sequences), aptamers (i.e., nucleic acids that bind non-nucleic target molecules) have been disclosed. See, e.g., Blackwell et al., *Science* (1990) 250:1104-1110; Blackwell et al., *Science* (1990) 250:1149-1152; Tuerk et al., *Science* (1990) 249:505-510; Joyce, *Gene* (1989) 82:83-87; and U.S. Pat. No. 5,840,867 entitled "Aptamer analogs specific for biomolecules".

**[0513]** As applied to aptamers, the term "binding" specifically excludes the "Watson-Crick"-type binding interactions (i.e., A:T and G:C base-pairing) traditionally associated with the DNA double helix. The term "aptamer" thus refers to a nucleic acid or a nucleic acid derivative that specifically binds to a target molecule, wherein the target molecule is either (i) not a nucleic acid, or (ii) a nucleic acid or structural element thereof that is bound through mechanisms other than duplex- or triplex-type base pairing. Such a molecule is called a "non-nucleic molecule" herein.

**[0514]** VIII.A. Structures of Nucleic Acids

**[0515]** "Nucleic acids," as used herein, refers to nucleic acids that are isolated a natural source; prepared in vitro, using techniques such as PCR amplification or chemical synthesis; prepared in vivo, e.g., via recombinant DNA technology; or by any appropriate method. Nucleic acids may be of any shape (linear, circular, etc.) or topology (single-stranded, double-stranded, supercoiled, etc.). The term "nucleic acids" also includes without limitation nucleic acid derivatives such as peptide nucleic acids (PNA's) and polypeptide-nucleic acid conjugates; nucleic acids having at least one chemically modified sugar residue, backbone, internucleotide linkage, base, nucleoside, or nucleotide analog; as well as nucleic acids having chemically modified 5' or 3' ends; and nucleic acids having two or more of such modifications. Not all linkages in a nucleic acid need to be identical.

**[0516]** Nucleic acids that are aptamers are often, but need not be, prepared as oligonucleotides. Oligonucleotides include without limitation RNA, DNA and mixed RNA-DNA molecules having sequences of lengths that have minimum lengths of 2, 4, 6, 8, 10, 11, 12, 13, 14, 15, 17, 18, 19, 20, 21, 22, 23, 24 or 25 nucleotides, and maximum lengths of about 100, 75, 50, 40, 25, 20 or 15 or more nucleotides, irrespectively. In general, a minimum of 6 nucleotides, preferably 10 nucleotides, more preferably 14 to 20 nucleotides, is necessary to effect specific binding.

**[0517]** In general, the oligonucleotides may be single-stranded (ss) or double-stranded (ds) DNA or RNA, or conjugates (e.g., RNA molecules having 5' and 3' DNA "clamps") or hybrids (e.g., RNA:DNA paired molecules), or derivatives (chemically modified forms thereof). However, single-stranded DNA is preferred, as DNA is often less labile than RNA. Similarly, chemical modifications that enhance

an aptamer's specificity or stability are preferred. VIII.B. Chemical Modifications of Nucleic Acids

**[0518]** Chemical modifications that may be incorporated into aptamers and other nucleic acids include, with neither limitation nor exclusivity, base modifications, sugar modifications, and backbone modifications.

**[0519]** Base modifications: The base residues in aptamers may be other than naturally occurring bases (e.g., A, G, C, T, U, 5MC, and the like). Derivatives of purines and pyrimidines are known in the art; an exemplary but not exhaustive list includes aziridinylcytosine, 4-acetylcytosine, 5-fluorouracil, 5-bromouracil, 5-carboxymethylaminomethyl-2-thiouracil, 5-carboxymethylaminomethyluracil, inosine, N6-isopentenyladenine, 1-methyladenine, 1-methylpseudouracil, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine (5MC), N6-methyladenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5-methoxyuracil, 2-methylthio-N-6-isopentenyladenine, uracil-5-oxyacetic acid methylester, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid, and 2,6-diaminopurine. In addition to nucleic acids that incorporate one or more of such base derivatives, nucleic acids having nucleotide residues that are devoid of a purine or a pyrimidine base may also be included in aptamers.

**[0520]** Sugar modifications: The sugar residues in aptamers may be other than conventional ribose and deoxyribose residues. By way of non-limiting example, substitution at the 2'-position of the furanose residue enhances nuclease stability. An exemplary, but not exhaustive list, of modified sugar residues includes 2' substituted sugars such as 2'-O-methyl-, 2'-O-alkyl-, 2'-O-allyl-, 2'-S-alkyl-, 2'-S-allyl-, 2'-fluoro-, 2'-halo-, or 2'-azido-ribose, carbocyclic sugar analogs, alpha-anomeric sugars, epimeric sugars such as arabinose, xyloses or lyxoses, pyranose sugars, furanose sugars, sedoheptuloses, acyclic analogs and abasic nucleoside analogs such as methyl riboside, ethyl riboside or propylriboside.

**[0521]** Backbone modifications: Chemically modified backbones include, by way of non-limiting example, phosphorothioates, chiral phosphorothioates, phosphorodithioates, phosphotriesters, aminoalkylphosphotriesters, methyl and other alkyl phosphonates including 3'-alkylene phosphonates and chiral phosphonates, phosphinates, phosphoramidates including 3'-amino phosphoramidate and aminoalkylphosphoramidates, thionophosphoramidates, thionoalkylphosphonates, thionoalkylphosphotriesters, and boranophosphates having normal 3'-5' linkages, 2'-5' linked analogs of these, and those having inverted polarity wherein the adjacent pairs of nucleoside units are linked 3'-5' to 5'-3' or 2'-5' to 5'-2'. Chemically modified backbones that do not contain a phosphorus atom have backbones that are formed by short chain alkyl or cycloalkyl internucleoside linkages, mixed heteroatom and alkyl or cycloalkyl internucleoside linkages, or one or more short chain heteroatomic or heterocyclic internucleoside linkages, including without limitation morpholino linkages; siloxane backbones; sulfide, sulfoxide and sulfone backbones; formacetyl and thioformacetyl backbones; methylene formacetyl and thioformacetyl backbones; alkene containing backbones; sulfamate

backbones; methyleneimino and methylenehydrazino backbones; sulfonate and sulfonamide backbones; and amide backbones.

**[0522]** VIII.C. Preparation and Identification of Aptamers

**[0523]** In general, techniques for identifying aptamers involve incubating a preselected non-nucleic target molecule with mixtures (2 to 50 members), pools (50 to 5,000 members) or libraries (50 or more members) of different nucleic acids that are potential aptamers under conditions that allow complexes of target molecules and aptamers to form. By "different nucleic acids" it is meant that the nucleotide sequence of each potential aptamer may be different from that of any other member, that is, the sequences of the potential aptamers are random with respect to each other. Randomness can be introduced in a variety of manners such as, e.g., mutagenesis, which can be carried out *in vivo* by exposing cells harboring a nucleic acid with mutagenic agents, *in vitro* by chemical treatment of a nucleic acid, or *in vitro* by biochemical replication (e.g., PCR) that is deliberately allowed to proceed under conditions that reduce fidelity of replication process; randomized chemical synthesis, i.e., by synthesizing a plurality of nucleic acids having a preselected sequence that, with regards to at least one position in the sequence, is random. By "random at a position in a preselected sequence" it is meant that a position in a sequence that is normally synthesized as, e.g., as close to 100% A as possible (e.g., 5'-C-T-T-A-G-T-3') is allowed to be randomly synthesized at that position (C-T-T-N-G-T, wherein N indicates a randomized position where, for example, the synthesizing reaction contains 25% each of A, T, C and G; or x % A, w % T, y % C and z % G, wherein  $x+w+y+z=100$ ). In later stages of the process, the sequences are increasingly less randomized and consensus sequences may appear; in any event, it is preferred to ultimately obtain an aptamer having a unique nucleotide sequence.

**[0524]** Aptamers and pools of aptamers are prepared, identified, characterized and/or purified by any appropriate technique, including those utilizing *in vitro* synthesis, recombinant DNA techniques, PCR amplification, and the like. After their formation, target:aptamer complexes are then separated from the uncomplexed members of the nucleic acid mixture, and the nucleic acids that can be prepared from the complexes are candidate aptamers (at early stages of the technique, the aptamers generally being a population of a multiplicity of nucleotide sequences having varying degrees of specificity for the target). The resulting aptamer (mixture or pool) is then substituted for the starting aptamer (library or pool) in repeated iterations of this series of steps. When a limited number (e.g., a pool or mixture, preferably a mixture with less than 100 members, more preferably less than 10 members, most preferably 1, of nucleic acids having satisfactory specificity is obtained, the aptamer is sequenced and characterized. Pure preparations of a given aptamer are generated by any appropriate technique (e.g., PCR amplification, *in vitro* chemical synthesis, and the like).

**[0525]** For example, Tuerk and Gold (Science (1990) 249:505-510) disclose the use of a procedure termed "systematic evolution of ligands by exponential enrichment" (SELEX). In this method, pools of nucleic acid molecules that are randomized at specific positions are subjected to selection for binding to a nucleic acid-binding protein (see,

e.g., PCT International Publication No. WO 91/19813 and U.S. Pat. No. 5,270,163). The oligonucleotides so obtained are sequenced and otherwise characterized. Kinzler, K. W., et al. (Nucleic Acids Res. (1989) 17:3645-3653) used a similar technique to identify synthetic double-stranded DNA molecules that are specifically bound by DNA-binding polypeptides. Ellington, A. D., et al. (Nature (1990) 346: 818-822) disclose the production of a large number of random sequence RNA molecules and the selection and identification of those that bind specifically to specific dyes such as Cibacron blue.

**[0526]** Another technique for identifying nucleic acids that bind non-nucleic target molecules is the oligonucleotide combinatorial technique disclosed by Ecker, D. J. et al. (Nuc. Acids Res. 21, 1853 (1993)) known as "synthetic unrandomization of randomized fragments" (SURF), which is based on repetitive synthesis and screening of increasingly simplified sets of oligonucleotide analogue libraries, pools and mixtures (Tuerk et al., Science 249:505, 1990). The starting library consists of oligonucleotide analogues of defined length with one position in each pool containing a known analogue and the remaining positions containing equimolar mixtures of all other analogues. With each round of synthesis and selection, the identity of at least one position of the oligomer is determined until the sequences of optimized nucleic acid ligand aptamers are discovered.

**[0527]** Once a particular candidate aptamer has been identified through a SURF, SELEX or any other technique, its nucleotide sequence can be determined (as is known in the art), and its three-dimensional molecular structure can be examined by nuclear magnetic resonance (NMR). These techniques are explained in relation to the determination of the three-dimensional structure of a nucleic acid ligand that binds thrombin in Padmanabhan et al., J. Biol. Chem. 24, 17651 (1993); Wang et al., Biochemistry 32, 1899 (1993); and Macaya et al., Proc. Nat'l. Acad. Sci. USA 90, 3745 (1993). Selected aptamers may be resynthesized using one or more modified bases, sugars or backbone linkages. Aptamers consist essentially of the minimum sequence of nucleic acid needed to confer binding specificity, but may be extended on the 5' end, the 3' end, or both, or may be otherwise derivatized or conjugated.

**[0528]** IX. Polypeptidic Binding Moieties

**[0529]** A variety of binding moieties can be attached to a minicell of the invention for a variety of purposes. In a preferred embodiment, the binding moiety is directed to a ligand that is displayed by a cell into which it is desired to deliver the therapeutic content of a minicell.

**[0530]** IX.A. Antibodies and Antibody Derivatives

**[0531]** The term "antibody" is meant to encompass an immunoglobulin molecule obtained by *in vitro* or *in vivo* generation of an immunogenic response, and includes polyclonal, monospecific and monoclonal antibodies, as well as antibody derivatives, e.g. single-chain antibody fragments (scFv). An "immunogenic response" is one that results in the production of antibodies directed to one or more proteins after the appropriate cells have been contacted with such proteins, or polypeptide derivatives thereof, in a manner such that one or more portions of the protein function as epitopes. An epitope is a single antigenic determinant in a molecule. In proteins, particularly denatured proteins, an

epitope is typically defined and represented by a contiguous amino acid sequence. However, in the case of nondenatured proteins, epitopes also include structures, such as active sites, that are formed by the three-dimensional folding of a protein in a manner such that amino acids from separate portions of the amino acid sequence of the protein are brought into close physical contact with each other.

**[0532]** Wildtype antibodies have four polypeptide chains, two identical heavy chains and two identical light chains. Both types of polypeptide chains have constant regions, which do not vary or vary minimally among antibodies of the same class (i.e., IgA, IgM, etc.), and variable regions. Variable regions are unique to a particular antibody and comprise an “antigen binding domain” that recognizes a specific epitope. Thus, an antibody’s specificity is determined by the variable regions located in the amino terminal regions of the light and heavy chains.

**[0533]** As used herein, the term “antibody” encompasses derivatives of antibodies such as antibody fragments that retain the ability to specifically bind to antigens. Such antibody fragments include Fab fragments (i.e., an antibody fragment that contains the antigen-binding domain and comprises a light chain and part of a heavy chain bridged by a disulfide bond); Fab’ (an antibody fragment containing a single anti-binding domain comprising an Fab and an additional portion of the heavy chain through the hinge region); F(ab’)<sub>2</sub> (two Fab’ molecules joined by interchain disulfide bonds in the hinge regions of the heavy chains; the Fab’ molecules may be directed toward the same or different epitopes); a bispecific Fab (an Fab molecule having two antigen binding domains, each of which may be directed to a different epitope); a single chain Fab chain comprising a variable region, a.k.a., a scFv (the variable, antigen-binding determinative region of a single light and heavy chain of an antibody linked together by a chain of about 10 to about 25 amino acids).

**[0534]** The term “antibody” includes antibodies and antibody derivatives that are produced by recombinant DNA techniques and “humanized” antibodies. Humanized antibodies have been modified, by genetic manipulation and/or in vitro treatment to be more human, in terms of amino acid sequence, glycosylation pattern, etc., in order to reduce the antigenicity of the antibody or antibody fragment in an animal to which the antibody is intended to be administered (Gussow et al., *Methods Enz.* 203:99-121, 1991).

**[0535]** A single-chain antibody (scfv) is a non-limiting example of a binding moiety that may be displayed on minicells. Single-chain antibodies are produced by recombinant DNA technology and may be incorporated into fusion proteins. The term “single chain” denotes the fact that scFv’s are found in a single polypeptide. In contrast, wildtype antibodies have four polypeptide chains, two identical heavy chains and two identical light chains. Both types of polypeptide chains have constant regions, which do not vary or vary minimally among antibodies of the same class (i.e., IgA, IgM, etc.), and variable regions. An antibody’s specificity is determined by the variable regions located in the amino terminal regions of the light and heavy chains. The variable regions of a light chain and associated heavy chain form an “antigen binding domain” that recognizes a specific epitope. In a single chain antibody, the amino acid sequences of the variable light and variable heavy regions of an antibody are

present in one contiguous polypeptide. Methods of producing single chain antibodies are known in the art. See, for example, U.S. Pat. Nos. 4,946,778; 5,260,203; 5,455,030; 5,518,889; 5,534,621; 5,869,620; 6,025,165; 6,027,725 and 6,121,424.

**[0536]** Antibody derivatives and other polypeptides that are binding moieties can be isolated from protein display libraries, in which a library of candidate binding agents is displayed on a phage or other agent that comprises a nucleic acid encoding the protein it displays. Thus, an agent that binds to the target compound can be isolated, and nucleic acid prepared therefrom, providing for the rapid isolation of binding moieties and nucleic acids that can be used to produce them. For reviews, see Benhar I. *Biotechnological applications of phage and cell display.* *Biotechnology Adv.* 2001 (19):1-33; FitzGerald K. *In vitro display technologies—new tools for drug discovery.* *Drug Discov Today.* 2000 5(6):253-258; and Hoogenboom H R, Chames P. *Natural and designer binding sites made by phage display technology.* *Immunol Today.* 2000 August;21(8):371-8.

**[0537]** A variety of protein display systems are known in the art and include various phage display systems such as those described in Jung S, Arndt K, Müller K, Plückthyn A. *Selectively infective phage (S1P) technology: scope and limitations.* *J Immunol Methods.* 1999 (231):93-104; Katz B. *Structural and mechanistic determinants of affinity and specificity of ligands discovered or engineered by phage display.* *Annu Rev Biophys Biomol Struct.* 1997 (26):27-45; Forrer P, Jung S, Pluckthun A. *Beyond binding: using phage display to select for structure, folding and enzymatic activity in proteins.* *Curr Opin Struct Biol.* 1999 August;9(4):514-20; Rondot S, Koch J, Breitling F, Dubel S. *A helper phage to improve single-chain antibody presentation in phage display.* *Nat Biotechnol.* 2001 January; 19(1):75-8. Giebel L B, Cass R T, Milligan D L, Young D C, Arze R, Johnson C R. *Screening of cyclic peptide phage libraries identifies ligands that bind streptavidin with high affinities.* *Biochemistry.* Nov. 28, 1995;34(47):15430-5; de Kruijff J, Logtenberg T. *Leucine zipper dimerized bivalent and bispecific scFv antibodies from a semi-synthetic antibody phage display library.* *J. Biol. Chem.* Mar. 29, 1996;271(13):7630-4; Hoogenboom H R, Henderikx P, de Haard H. *Creating and engineering human antibodies for immunotherapy.* *Adv Drug Deliv Rev.* Apr. 6, 1998;31(1-2):5-31; Helfrich W, Haisma H J, Magdolen V, Luther T, Bom V J, Westra J, van der Hoeven R, Kroesen B J, Molema G, de Leij L. *A rapid and versatile method for harnessing scFv antibody fragments with various biological effector functions.* *J Immunol Methods.* Apr. 3, 2000;237(1-2):131-45; Hoess R H. *Bacteriophage lambda as a vehicle for peptide and protein display.* *Curr Pharm Biotechnol* 2002 March;3(1):23-8; Baek H, Suk K H, Kim Y H, Cha S. *An improved helper phage system for efficient isolation of specific antibody molecules in phage display.* *Nucleic Acids Res.* Mar. 1, 2002;30(5):e18; and Rondot S, Koch J, Breitling F, Dubel S. *A helper phage to improve single-chain antibody presentation in phage display.* *Nat Biotechnol.* 2001 January;19(1):75-8.

**[0538]** Other display systems include without limitation “Yeast Display” (*Curr Opin Biotechnol* 1999 October; 10(5):422-7. *Applications of yeast in biotechnology: protein production and genetic analysis.* Cereghino G P, Cregg J M.); “Baculovirus Display” (Kost T A, Condreay J P.

- Recombinant baculoviruses as expression vectors for insect and mammalian cells. *Curr Opin Biotechnol.* 1999 October; 10(5):428-33; and Liang M, Dubel S, Li D, Queitsch I, Li W, Bautz E K. Baculovirus expression cassette vectors for rapid production of complete human IgG from phage display selected antibody fragments. *J Immunol Methods.* Jan. 1, 2001;247(1-2):119-30; "Ribosome Display" (Hanes J, Schaffitzel C, Knappik A, Pluckthun A. Picomolar affinity antibodies from a fully synthetic naive library selected and evolved by ribosome display. *Nat Biotechnol.* 2000 December;18(12):1287-92; Hanes J, Jermutus L, Pluckthun A. Selecting and evolving functional proteins in vitro by ribosome display. *Methods Enzymol.* 2000;328:404-30; Schaffitzel C, Hanes J, Jermutus L, Pluckthun A. Ribosome display: an in vitro method for selection and evolution of antibodies from libraries. *J Immunol Methods.* Dec. 10, 1999;231(1-2):119-35; Hanes J, Jermutus L, Weber-Bornhauser S, Bosshard H R, Pluckthun A. Ribosome display efficiently selects and evolves high-affinity antibodies in vitro from immune libraries. *Proc Natl Acad Sci USA.* Nov. 24, 1998;95(24): 14130-5; Hanes J, Pluckthun A. In vitro selection and evolution of functional proteins by using ribosome display. *Proc Natl Acad Sci U S A.* May 13, 1997;94(10):4937-42; Coia G, Pontes-Braz L, Nuttall S D, Hudson P J, Irving R A. Panning and selection of proteins using ribosome display. *J Immunol Methods.* Aug. 1, 2001;254(1-2):191-7.; Irving R A, Coia G, Roberts A, Nuttall S D, Hudson P J. Ribosome display and affinity maturation: from antibodies to single V-domains and steps towards cancer therapeutics. *J Immunol Methods.* Feb. 1, 2001;248(1-2):31-45); and "Bacterial Display" (Hoischen C, Fritsche C, Gumpert J, Westermann M, Gura K, Fahnert B. Novel bacterial membrane surface display system using cell wall-less L-forms of *Proteus mirabilis* and *Escherichia coli*. *Appl Environ Microbiol.* 2002 February;68(2):525-31; Etz H, Minh D B, Schellack C, Nagy E, Meinke A. Bacterial phage receptors, versatile tools for display of polypeptides on the cell surface. *J Bacteriol.* 2001 December;183(23):6924-35; Patel D, Vitovski S, Senior H J, Edge M D, Hockney R C, Dempsey M J, Sayers J R. Continuous affinity-based selection: rapid screening and simultaneous amplification of bacterial surface-display libraries. *Biochem J.* Aug. 1, 2001;357(Pt 3):779-85; Lang H. Outer membrane proteins as surface display systems. *Int J Med Microbiol.* 2000 December;290(7):579-85; Earhart C F. Use of an Lpp-OmpA fusion vehicle for bacterial surface display. *Methods Enzymol.* 2000;326:506-16; Benhar I, Azriel R, Nahary L, Shaky S, Berdichevsky Y, Tamarkin A, Wels W. Highly efficient selection of phage antibodies mediated by display of antigen as Lpp-OmpA fusions on live bacteria. *J. Mol. Biol.* Aug. 25, 2000;301(4):893-904; Xu Z, Lee S Y. Display of polyhistidine peptides on the *Escherichia coli* cell surface by using outer membrane protein C as an anchoring motif. *Appl Environ Microbiol.* 1999 November;65(11):5142-7; Daugherty P S, Olsen M J, Iverson B L, Georgiou G. Development of an optimized expression system for the screening of antibody libraries displayed on the *Escherichia coli* surface. *Protein Eng.* 1999 July;12(7):613-21; Chang H J, Sheu S Y, Lo S J. Expression of foreign antigens on the surface of *Escherichia coli* by fusion to the outer membrane protein traT. *J Biomed Sci.* 1999 January;6(1):64-70; Maurer J, Jose J, Meyer T F. Autodisplay: one-component system for efficient surface display and release of soluble recombinant proteins from *Escherichia coli*. *J Bacteriol.* 1997 February; 179(3):794-804.
- [0539] Antibodies, particularly single-chain antibodies, directed to surface antigens specific for a particular cell type may also be used as cell- or tissue-specific targeting elements. Single-chain antibody amino acid sequences have been incorporated into a variety of fusion proteins, including those with transmembrane domains and/or membrane-anchoring domains. See, for example, Kuroki et al., "Specific Targeting Strategies of Cancer Gene Therapy Using a Single-Chain Variable Fragment (scFv) with a High Affinity for CEA," *Anticancer Res.*, pp. 4067-71, 2000; U.S. Pat. No. 6,146,885, to Dornburg, entitled "Cell-Type Specific Gene Transfer Using Retroviral Vectors Containing Antibody-Envelope Fusion Proteins"; Jiang et al., "In Vivo Cell Type-Specific Gene Delivery With Retroviral Vectors That Display Single Chain Antibodies," *Gene Ther.* 1999, 6:1982-7; Engelstadter et al., "Targeting Human T Cells By Retroviral Vectors Displaying Antibody Domains Selected From A Phage Display Library," *Hum. Gene Ther.* 2000, 11:293-303; Jiang et al., "Cell-Type-Specific Gene Transfer Into Human Cells With Retroviral Vectors That Display Single-Chain Antibodies," *J. Virol* 1998,72:10148-56; Chu et al., "Toward Highly Efficient Cell-Type-Specific Gene Transfer With Retroviral Vectors Displaying Single-Chain Antibodies," *J. Virol* 1997, 71:720-5; Chu et al., "Retroviral Vector Particles Displaying The Antigen-Binding Site Of An Antibody Enable Cell-Type-Specific Gene Transfer," *J. Virol* 1995, 69:2659-63; Chu et al., "Cell Targeting With Retroviral Vector Particles Containing Antibody-Envelope Fusion Proteins," *Gene Ther.* 1994, 1:292-9; Esshar et al., "Specific activation and targeting of cytotoxic lymphocytes through chimeric single chains consisting of antibody-binding domains and the or subunits of the immunoglobulin and T-cell receptors," *Proc. Natl. Acad. Sci. USA*, 1993, Vol. 90:720-724; Einfeld et al., "Construction of a Pseudoreceptor That Mediates Transduction by Adenoviruses Expressing a Ligand in Fiber or Penton Base," *J. Virol.* 1999, 73:9130-9136; Marin et al., "Targeted Infection of Human Cells via Major Histocompatibility Complex Class I Molecules by Moloney Murine Leukemia Virus-Derived Viruses Displaying Single-Chain Antibody Fragment-Envelope Fusion Proteins," *J. Virol.*, 1996, 70:2957-2962; Somia et al., "Generation of targeted retroviral vectors by using single-chain variable fragment: An approach to in vivo gene delivery," *Proc. Natl. Acad. Sci. USA*, 1995, 92:7570-7574; Liu et al., "Treatment of B-Cell Lymphoma With Chimeric IgG and Single-Chain Fv Antibody-Interleukin-2 Fusion Proteins," *Blood*, 1998, 92:2103-2112; Martin et al., "Retrovirus Targeting by Tropism Restriction to Melanoma Cells," *J. Virol.*, 1999, 73:6923-6929; Ramjiawan et al., "Noninvasive Localization of Tumors by Immunofluorescence Imaging Using a Single Chain Fv Fragment of a Human Monoclonal Antibody with Broad Cancer Specificity," *Amer. Cancer Society*, 2000, 89:1134-1144; Snitkovsky et al., "A TVA-Single-Chain Antibody Fusion Protein Mediates Specific Targeting of a Subgroup A Avian Leukosis Virus Vector to Cells Expressing a Tumor-Specific Form of Epidermal Growth Factor Receptor," *J. Virol.*, 2000, 74:9540-9545; Chu et al., "Toward Highly Efficient Cell-Type-Specific Gene Transfer with Retroviral Vectors Displaying Single-Chain Antibodies," *J. Virol.*, 1997, 71:720-725; Kulkarni et al., Pro-

grammed cell death signaling via cell-surface expression of a single-chain antibody transgene, *Transplantation* Mar. 27, 2000;69(6): 1209-17.

**[0540]** IX.B. Non-Catalytic Derivatives of Active Sites of Enzymes

**[0541]** Enzymes bind their substrates, at least transiently, in regions known as "active sites." It is known in the art that non-catalytic derivatives of enzymes, which bind but do not chemically alter their substrates may be prepared. Non-catalytic enzymes, particularly the mutant active sites thereof, are used to bind substrate molecules.

**[0542]** As a non-limiting example, enzymes from which biologically inactive (non-catalytic) sphingolipid-binding derivatives are obtained. Such derivatives of these enzymes bind their substrate sphingolipid. Sphingosine-1-phosphate (S1P) is bound by non-catalytic derivatives of enzymes having S1P as a substrate, e.g., S1P lyase and S1P phosphatase. Sphingosine (SPH) is bound by non-catalytic derivatives of enzymes having SPH as a substrate, e.g., SPH kinase and ceramide synthase. Ceramide (CER) is bound by non-catalytic derivatives of enzymes having CER as a substrate, such as, by way of non-limiting example, ceramidase, sphingomyelin synthase, ceramide kinase, and glucosylceramide synthase. Sphingomyelin is bound by non-catalytic derivatives of sphingomyelinase, an enzyme having sphingomyelin as a substrate.

**[0543]** IX.C. Nucleic Acid Binding Domains

**[0544]** Nucleic acid binding polypeptide domains may bind nucleic acids in a sequence-dependent or sequence-independent fashion and/or in a manner that is specific for various nucleic acids having different chemical structures (e.g., single- or double-stranded DNA or RNA, RNA:DNA hybrid molecules, etc.). Non-limiting examples of membrane-based transcription factors and DNA-binding protein include Smad proteins (Miyazono et al., TGF-beta signaling by Smad proteins (Review), *Adv Immunol* 75:115-57, 2000); SREBPs (sterol regulatory element binding proteins) (Ye et al., Asparagine-proline sequence within membrane-spanning segment of SREBP triggers intramembrane cleavage by site-2 protease, *Proc Natl Acad Sci USA* 97:5123-8, 2000; Shimomura et al., Cholesterol feeding reduces nuclear forms of sterol regulatory element binding proteins in hamster liver, *Proc Natl Acad Sci USA* 94:12354-9, 1997; Brown and Goldstein, The SREBP pathway: regulation of cholesterol metabolism by proteolysis of a membrane-bound transcription factor (Review), *Cell* 89:331-40, 1997; Scheek et al., Sphingomyelin depletion in cultured cells blocks proteolysis of sterol regulatory element binding proteins at site 1, *Proc Natl Acad Sci USA* 94:11179-83, 1997); mitochondrial DNA-binding membrane proteins, e.g., Abf2p and YhmZp (Cho et al., A novel DNA-binding protein bound to the mitochondrial inner membrane restores the null mutation of mitochondrial histone Abf2p in *Saccharomyces cerevisiae*, *Mol Cell Biol* 18:5712-23, 1998); and bacterial DNA-binding membrane proteins (Smith et al., Transformation in *Bacillus subtilis*: purification and partial characterization of a membrane-bound DNA-binding protein., *J Bacteriol* 156:101-8, 1983).

**[0545]** IX.D. Attaching Binding Moieties, or Other Compounds, to Minicells

**[0546]** Binding compounds-or moieties can be chemically attached (conjugated) to minicells via membrane proteins

that are displayed on the minicells. The compound to be conjugated to minicells (the "attachable compound") may of any chemical composition, i.e., a small molecule, a nucleic acid, a radioisotope, a lipid or a polypeptide. One type of attachable compound that can be covalently attached to minicells is a binding moiety, e.g., an antibody or antibody derivative. Another non-limiting example of attachable compounds is polyethylene glycol ("PEG"), which lowers the uptake *in vivo* of minicells by the reticuloendothelial system (RES). Another non-limiting example of creating stealth minicells to avoid the RES is to express proteins or other molecules on the surfaces of minicells whose lipid compositions have been modified, such as anionic lipid-rich minicells.

**[0547]** By way of non-limiting example, it is possible to prepare minicells that express transmembrane proteins with cysteine moieties on extracellular domains. Linkage of the membrane protein may be achieved through surface cysteinyl groups by, e.g., reduction with cysteinyl residues on other compounds to form disulfide bridges (S=S). If appropriate cysteinyl residues are not present on the membrane protein they may be introduced by genetic manipulation. The substitution of cysteine for another amino acid may be achieved by methods well-known to those skilled in the art, for example, by using methods described in Maniatis, Sambrook, and Fritsch (*Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, 1989). As a non-limiting example, bioactive lysosphingolipids (e.g., sphingosine, sphingosine-1-phosphate, sphingosylphosphoryl choline) are covalently linked to proteins expressed on the surfaces of minicells such that these bioactive lipids are on the surface of the minicells and accessible for therapeutic or diagnostic uses *in vivo* or *in vitro*.

**[0548]** When the attachable moiety and the membrane protein both have a reduced sulfhydryl group, a homobifunctional cross-linker that contains maleimide, pyridyl disulfide, or beta-alpha-haloacetyl groups may be used for cross-linking. Examples of such cross-linking reagents include, but are not limited to, bismaleimidohexane (BMH) or 1,4-Di-[3'-(2'-pyridyldithio)propionamido]butane (DPDPB). Alternatively, a heterobifunctional cross-linker that contains a combination of maleimide, pyridyl disulfide, or beta-alpha-haloacetyl groups can be used for cross-linking.

**[0549]** As another non-limiting example, attachable moieties may be chemically conjugated using primary amines. In these instances, a homobifunctional cross-linker that contains succinimide ester, imidoester, acylazide, or isocyanate groups may be used for cross-linking. Examples of such cross-linking reagents include, but are not limited to: Bis[2-(succinimidooxycarbonyloxy)ethyl]sulfone (BSO-COES); Bis[2-(sulfosuccinimidooxycarbonyloxy)ethyl]sulfone (sulfo-BSCOCOES); Disuccinimidyl suberate (DSS); Bis-(Sulfosuccinimidyl) Suberate (BS3); Disuccinimidyl glutarate (DSG); Dithiobis(succinimidylpropionate) (DSP); Dithiobis(sulfosuccinimidylpropionate) (DTSSP); Disulfosuccinimidyl tartrate (sulfo-DST); Dithio-bis-maleimidodethane (DTME); Disuccinimidyl tartrate (DST); Ethylene glycolbis(sulfosuccinimidylsuccinate) (sulfo-EGS); Dimethyl malonimidate.2 HCl (DMM); Ethylene glycolbis(succinimidylsuccinate) (EGS); Dimethyl succinimidate.2 HCl (DMSC); Dimethyl adipimidate.2 HCl (DMA); Dimethyl pimelimidate.2 HCl (DMP); and Dimethyl



suberimidate.2.HCl (DMS), and Dimethyl 3,3'-dithiobispropionimidate.2 HCl (DTBP). Heterobifunctional cross-linkers that contains a combination of imidoester or succinimide ester groups may also be used for cross-linking.

[0550] As another non-limiting example, attachable moieties may be chemically conjugated using sulfhydryl and primary amine groups. In these instances, heterobifunctional cross-linking reagents are preferable used. Examples of such cross-linking reagents include, but are not limited to: N-succinimidyl 3-(2-pyridyldithio)propionate (DPDP); N-succinimidyl 6-[3'-(2-pyridyldithio)-propionamido] hexanoate (sulfo-LC-SPDP); m-maleimidobenzoyl-N-hydroxysuccinimide ester (MBS); m-maleimidobenzoyl-N-hydroxysulfosuccinimide ester (sulfo-MBS); succinimidyl 4-[p-maleimidophenyl] butyrate (SMPB); sulfosuccinimidyl 4-[p-maleimidophenyl] butyrate (sulfo-SMPB); N-[γ-Maleimidobutyryloxy] succinimide ester (GMBS), N-[γ-maleimidobutyryloxy] sulfosuccinimide ester (sulfo-GMBS); N-[ε-maleimidocaproyloxy] succinimide ester (EMCS); N-[ε-maleimidocaproyloxy] sulfosuccinimide ester (sulfo-EMCS); N-succinimidyl(4-iodoacetyl)aminobenzoate (SIAB); sulfosuccinimidyl(4-iodoacetyl)aminobenzoate (sulfo-SIAB); succinimidyl 4-(N-maleimidomethyl)cyclohexane-1-carboxylate (SMCC); sulfosuccinimidyl 4-(N-maleimidomethyl)cyclohexane-1-carboxylate (sulfo-SMCC); succinimidyl-4-(N-maleimidomethyl)cyclohexane-1-carboxylate (6-amido-caproate) (LC-SMCC); 4-succinimidylloxycarbonyl-methyl-(2-pyridyldithio) toluene (SMPT); and sulfo-LC-SMPT.

[0551] As an exemplary protocol, a minicell suspension is made 5 mM EDTA/PBS, and a reducing solution of 2-mercaptoethylamine in 5 mM EDTA/PBS is added to the minicells. The mixture is incubated for 90 minutes at 37° C. The minicells are washed with EDTA/PBS to remove excess 2-mercaptoethylamine. The attachable moiety is dissolved in PBS, pH 7.2. A maleimide crosslinker is added to the solution, which is then incubated for 1 hour at room temperature. Excess maleimide is removed by column chromatography.

[0552] The minicells with reduced sulfhydryl groups are mixed with the derivatized compounds having an attachable moiety. The mixture is allowed to incubate at 4° C. for 2 hours or overnight to allow maximum coupling. The conjugated minicells are washed to remove unreacted (unattached) compounds having the attachable moiety. Similar protocols are used for expressed membrane proteins with other reactive groups (e.g., carboxyl, amine) that can be conjugated to an attachable moiety.

[0553] IX.E. Non-Genetic Methods for Directing Compounds to Membranes

[0554] Included within the scope of the invention are compounds that can be inserted into the membrane of segregated minicells. Such compounds include attachable moieties that are chemically conjugated to the surface of a minicell, and compounds that associate with and/or insert into a membrane "spontaneously," i.e., by virtue of their chemical nature. By way of non-limiting example, proteins that "spontaneously" insert into membranes include but are not limited to Thykaloid membrane proteins (Woolhead et al., *J. Biol. Chem.* 276:14607-14613, 2001), the mitochondrial adenine nucleotide translocator (Jacotot et al., *J. Exp. Med.* 193:509-519, 2001), and polypeptides obtained using

the methods of Hunt et al. (Spontaneous, pH-dependent membrane insertion of a transbilayer alpha-helix, *Biochem* 36:15177-15192, 1997). Lipids, gangliosides, sphingomyelins, plasmalogens glycosyl diacylglycerols, and sterols can also be incorporated into the membranes of segregated minicells.

[0555] X. Membrane Proteins

[0556] In certain aspects of the invention, membrane proteins from non-eubacterial organisms are expressed and displayed by minicells. The cellular membrane (a.k.a. the "plasma membrane") is a lipid bilayer that forms the boundary between the interior of a cell and its external environment. The term "membrane proteins" refers to proteins that are found in membranes including without limitation cellular and organellar membranes.

[0557] X.A. Types of Membrane Proteins

[0558] X.A.1. In General

[0559] Membrane proteins consist, in general, of two types, peripheral membrane proteins and integral membrane proteins.

[0560] Integral membrane proteins can span both layers (or "leaflets") of a lipid bilayer. Thus, such proteins may have extracellular, transmembrane, and intracellular domains. Extracellular domains are exposed to the external environment of the cell, whereas intracellular domains face the cytosol of the cell. The portion of an integral membrane protein that traverses the membrane is the "transmembrane domain." Transmembrane domains traverse the cell membrane often by one or more regions comprising 15 to 25 hydrophobic amino acids which are predicted to adopt an alpha-helical conformation.

[0561] Integral membrane proteins are classified as bitopic or polytopic (Singer, (1990) *Annu. Rev. Cell Biol.* 6:247-96). Bitopic proteins span the membrane once while polytopic proteins contain multiple membrane-spanning segments.

[0562] A peripheral membrane protein is a membrane protein that is bound to the surface of the membrane and is not integrated into the hydrophobic layer of a membrane region. Peripheral membrane proteins do not span the membrane but instead are bound to the surface of a membrane, one layer of the lipid bilayer that forms a membrane, or the extracellular domain of an integral membrane protein.

[0563] X.A.2. In General

[0564] The invention can be applied to any membrane protein, including but not limited to the following exemplary receptors and membrane proteins. The proteins include but are not limited to are receptors (e.g., GPCRs, sphingolipid receptors, neurotransmitter receptors, sensory receptors, growth factor receptors, hormone receptors, chemokine receptors, cytokine receptors, immunological receptors, and compliment receptors, FC receptors), channels (e.g., potassium channels, sodium channels, calcium channels.), pores (e.g., nuclear pore proteins, water channels), ion and other pumps (e.g., calcium pumps, proton pumps), exchangers (e.g., sodium/potassium exchangers, sodium/hydrogen exchangers, potassium/hydrogen exchangers), electron transport proteins (e.g., cytochrome oxidase), enzymes and kinases (e.g., protein kinases, ATPases, GTPases, phos-

phatases, proteases.), structural/linker proteins (e.g., Caveolins, clathrin), adapter proteins (e.g., TRAD, TRAP, FAN), chemotactic/adhesion proteins (e.g., ICAM11, selectins, CD34, VCAM-1, LFA-1, VLA-1), and phospholipases such as PI-specific PLC and other phospholipases.

[0565] X.A.3. Receptors

[0566] Within the scope of the invention are any receptor, including without limitation:

[0567] The nuclear receptors, e.g the nuclear export receptor;

[0568] The peripheral (mitochondrial) benzodiazepine receptor (Gavish et al., "Enigma of the Peripheral Benzodiazepine Receptor," *Pharmacological Reviews*, Vol. 51, No. 4);

[0569] Adrenergic and muscarinic receptors (Brodde et al., "Adrenergic and Muscarinic Receptors in the Human Heart", *Pharmacological Review*, Vol. 51, No. 4);

[0570] Gamma-aminobutyric acid<sub>A</sub> receptors (Barnard et al., "International Union of Pharmacology. IV. Subtypes of  $\gamma$ -Aminobutyric Acid<sub>A</sub> Receptors: Classification on the Basis of Submit Structure and Receptor Function," *Pharmacological Reviews*, Vol. 50, No. 2);

[0571] Kinin Bi receptors (Marceau et al., "The Bi Receptors for Kinins," *Pharmacological Reviews*, Vol. 50, No. 3);

[0572] Chemokine receptors (Murphy et al., "International Union of Pharmacology. XXII. Nomenclature for Chemokine Receptors" *Pharmacological Review*, Vol. 52, No. 1);

[0573] Glycine and NMDA Receptors (Danysz et al., "Glycine and N-Methyl-D-Aspartate Receptors: Physiological Significance and Possible Therapeutic Applications," *Pharmacological Reviews*, Vol. 50, No. 4);

[0574] Glutamate receptor ion channels (Dingledine et al., "The Glutamate Receptor Ion Channels", *Pharmacological Reviews*, Vol. 51, No. 1);

[0575] Purine and pyrimidine receptors including purinergic (e.g., P2) receptors (Ralevic et al., "Receptors for Purines and Pyrimidines", *Pharmacological Reviews*, Vol. 50, No. 3); CNS receptors and membrane transporters (E. Sylvester Vizi, "Role of High-Affinity Receptors and Membrane Transporters in Nonsynaptic Communication and Drug Action in the Central Nervous System," *Pharmacological Reviews*, Vol. 52, No. 1);

[0576] Opioid receptors, including but not limited to the  $\mu$ -opioid receptor (Quock et al., "The  $\mu$ -Opioid Receptor: Molecular Pharmacology, Signal Transduction and the Determination of Drug Efficacy", *Pharmacological Review*, Vol. 51, No. 3);

[0577] Angiotensin II receptors (Gasparo et al., "International Union of Pharmacology. XXIII. The Angiotensin II Receptors" *Pharmacological Review*, Vol. 52, No. 3);

[0578] Cholecystokinin receptors (Noble et al., "International Union of Pharmacology. XXI. Structure, Distribution, and Functions of Cholecystokinin Receptors", *Pharmacological Reviews*, Vol. 51, No. 4)

[0579] Hormone receptors, including but not limited to, the estrogen receptor; the glucocorticoid receptor; and the insulin receptor;

[0580] Receptors found predominantly in the central nervous system, including but not limited to, neuronal nicotinic acetylcholine receptors; the dopamine D2/D3 receptor; GABA receptors; central cannabinoid receptor CB1; opioid receptors, e.g., the kappa opioid receptor, and the methadone-specific opioid receptor; nicotinic acetylcholine receptors; serotonin receptors, e.g., the serotonin 5-HT<sub>3</sub> receptor, the serotonin 5-HT<sub>4</sub> receptor, and the serotonin-2 receptor; and dopamine receptors, e.g., the dopamine D2/D3 receptor; and the neurotensin receptor;

[0581] Receptors for growth factors, including but not limited to, the erythropoietin receptor; the FGF receptor; the EGF receptor; the VEGF receptor; VEGF receptor-2 protein; VEGF-receptor protein (KDR); fibroblast growth factor receptor; the p75 nerve growth factor receptor; epidermal growth factor receptor; IGF-1 receptor; platelet factor-4 receptor; alpha platelet-derived growth factor receptor; hepatocyte growth factor receptor; and human fibroblast growth factor receptor;

[0582] Receptors for sphingolipids and lysophospholipids such as the Edg family of GPCRs;

[0583] Receptors for interleukins, e.g., receptors for interleukin-1 (IL-1), IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, et seq.; and

[0584] Various receptors, including by way of non-limiting example, receptors described in U.S. Pat. No. 6,210,967 (DNA encoding a mammalian LPA receptor and uses thereof); U.S. Pat. No. 6,210,921 (CAR: a novel coxsackievirus and adenovirus receptor; U.S. Pat. No. 6,211,343 (Lactoferrin receptor protein; U.S. Pat. No. 6,218,509 (LH/CG receptor, DNA and use thereof; U.S. Pat. No. 6,214,972 (DNA encoding prostaglandin receptor DP); U.S. Pat. No. 6,221,613 (DNA encoding a human melanin concentrating hormone receptor (MCH1) and uses thereof); U.S. Pat. No. 6,221,660 (DNA encoding SNORF25 receptor); U.S. Pat. No. 6,225,080 ( $\mu$ -subtype opioid receptor); U.S. Pat. No. 6,222,015 (Estrogen receptor); U.S. Pat. No. 6,228,610 (Human metabotropic glutamate receptor subtypes (hmR4, hmR6, hmR7) and related DNA compounds); U.S. Pat. No. 6,235,496 (Nucleic acid encoding mammalian  $\mu$  opioid receptor); U.S. Pat. No. 6,258,556 (cDNA and genomic clones encoding human  $\mu$  opiate receptor and the purified gene product); U.S. Pat. No. 6,245,531 (Polynucleotide encoding insect ecdysone receptor); U.S. Pat. No. 6,225,531 (Glucan elicitor receptor, DNA molecule coding therefor, fungus-resistant plants transformed with the DNA molecule and method for creating the plants); U.S. Pat. No. 6,245,893 (Receptor that binds anti-convulsant compounds); U.S. Pat. No. 6,248,712 (Urokinase-type plasminogen activator receptor; U.S. Pat. No. 6,248,554 (DNA sequence coding for a BMP receptor); U.S. Pat. No. 6,248,520 (Nucleic acid molecules encoding nuclear hormone receptor coactivators and uses thereof); U.S. Pat. No. 6,242,251 (Rhesus neuropeptide Y5 receptor); U.S. Pat. No. 6,252,056 (Human lysophosphatidic acid receptor and use thereof); U.S. Pat. No. 6,255,472 (Isolated nucleic acid molecule encoding a human skeletal muscle-specific receptor); U.S. Pat. No. 6,291,207 (Herpes virus entry receptor protein); U.S. Pat. No. 6,291,206 (BMP receptor proteins); U.S. Pat. No. 6,291,195 (DNA encoding a human melanin concentrating hormone receptor (MCH1) and uses thereof); U.S. Pat. No. 6,344,200 (Lactoferrin receptor protein); U.S. Pat. No.

6,335,180 (Nucleic acid sequences encoding capsaicin receptor and uses thereof); U.S. Pat. No. 6,265,184 (Polynucleotides encoding chemokine receptor 88C); U.S. Pat. No. 6,207,799 (Neuropeptide Y receptor Y5 and nucleic acid sequences); U.S. Pat. No. 6,290,970 (Transferrin receptor protein of *Moraxella*); U.S. Pat. No. 6,326,350 (Transferrin receptor subunit proteins of *Neisseria meningitidis*); U.S. Pat. No. 6,313,279 (Human glutamate receptor and related DNA compounds); U.S. Pat. No. 6,313,276 (Human endothelin receptor); U.S. Pat. No. 6,307,030 (Androgen receptor proteins, recombinant DNA molecules coding for such, and use of such compositions); U.S. Pat. No. 6,306,622 (cDNA encoding a BMP type II receptor); U.S. Pat. No. 6,300,087 (DNA encoding a human serotonin receptor (5-HT4B) and uses thereof); U.S. Pat. No. 6,297,026 (Nucleic acids encoding the C140 receptor); U.S. Pat. No. 6,277,976 (Or-1, an orphan receptor belonging to the nuclear receptor family); U.S. Pat. No. 6,274,708 (Mouse interleukin-11 receptor); U.S. Pat. No. 6,271,347 (Eosinophil eotaxin receptor); U.S. Pat. No. 6,262,016 (Transferrin receptor genes); U.S. Pat. No. 6,261,838 (Rat melanocortin receptor MC3-R); U.S. Pat. No. 6,258,943 (Human neurokinin-3 receptor); U.S. Pat. No. 6,284,870 (Gamma retinoic acid receptor); U.S. Pat. No. 6,258,944 (OB receptor isoforms and nucleic acids encoding them); 6,261,801 (Nucleic acids encoding tumor necrosis factor receptor 5); U.S. Pat. No. 6,261,800 (Luteinizing hormone/choriogonadotropin (LH/CG) receptor); U.S. Pat. No. 6,265,563 (Opioid receptor genes); U.S. Pat. No. 6,268,477 (Chemokine receptor 88-C); U.S. Pat. No. 6,316,611 (Human N-methyl-D-aspartate receptor subunits, nucleic acids encoding same and uses thereof); U.S. Pat. No. 6,316,604 (Human C3b/C4b receptor (CRI)); U.S. Pat. No. 6,287,855 (Nucleic acid encoding rat galanin receptor (GALR2)); U.S. Pat. No. 6,268,221 (Melanocyte stimulating hormone receptor and uses); and U.S. Pat. No. 6,268,214 (Vectors encoding a modified low affinity nerve growth factor receptor).

#### [0585] X.A.3. Other Membrane Proteins

[0586] Other membrane proteins are within the scope of the invention and include but are not limited to channels (e.g., potassium channels, sodium channels, calcium channels), pores (e.g., nuclear pore proteins, water channels), ion and other pumps (e.g., calcium pumps, proton pumps), exchangers (e.g., sodium/potassium exchangers, sodium/hydrogen exchangers, potassium/hydrogen exchangers), electron transport proteins (e.g., cytochrome oxidase), enzymes and kinases (e.g., protein kinases, ATPases, GTPases, phosphatases, proteases.), structural/linker proteins (e.g., Caveolins, clathrin), adapter proteins (e.g., TRAD, TRAP, FAN),

#### [0587] X.A.3.a. Cellular Adhesion Molecules

[0588] Cellular adhesion molecules, including but not limited to human rhinovirus receptor (ICAM-1), ICAM-2, ICAM-3, and PECAM-1, and chemotactic/adhesion proteins (e.g., selectins, CD34, VCAM-1, LFA-1, VLA-1) are within the scope of the invention. See also Alpin et al., "Signal Transduction and Signal Modulation by Cell Adhesion Receptors: The Role of Integrins, Cadherins, Immunoglobulin-Cell Adhesion Molecules, and Selectins", *Pharmacological Reviews*, Vol. 50, No. 2.

#### [0589] X.A.3.b. Cytochrome P450 Enzymes

[0590] The family of enzymes known as "cytochrome P450" enzymes (since they absorb light in the 450 nanom-

eter range), or as "cytochrome oxidase" enzymes (since they oxidize a wide range of compounds that do not naturally occur in circulating blood), are included within the scope of the invention. P450 enzymes encompasses a variety of enzymes, many of which are involved in xenobiotic metabolism, including by way of non-limiting example the metabolism of drugs, prodrugs and toxins. Directories and databases of P450s, and information regarding their substrates, are available on-line (Fabian et al., *The Directory of P450-containing Systems* in 1996, *Nucleic Acids Research* 25:274-277, 1997). In humans, at least about 200 different P450s are present (for a review, see Hasler et al., *Human cytochromes P450*, *Molecular Aspects of Medicine* 20:1-137, 1999). There are multiple forms of these P450s and each of the individual forms exhibit degrees of specificity towards individual compounds or sets of compounds. In some cases, a substrate, whether it is a drug or a carcinogen, is metabolized by more than one cytochrome P450.

[0591] Members of the cytochrome P450 family are present in varying levels and their expression and activities are controlled by variables such as chemical environment, sex, developmental stage, nutrition and age. The cytochrome P450s are found at high concentrations in liver cells, and at lower concentrations in other organs and tissues such as the lungs (e.g., Forme-Pfister et al., *Xenobiotic and endobiotic inhibitors of cytochrome P-450 dhl function, the target of the debrisoquine/sparteine type polymorphism*, *Biochem. Pharmacol.* 37:3829-35, 1988). By oxidizing lipophilic compounds, which makes them more water-soluble, cytochrome oxidase enzymes help the body eliminate (via urine, or in aerosols exhaled out of the lungs) compounds that might otherwise act as toxins or accumulate to undesired levels.

[0592] In humans, several cytochrome P450s have been identified as being involved in xenobiotic metabolism. These include CYP1A1, CYP1A2, CYP2A6, CYP2B6, CYP2C8, CYP2C9, CYP2C18, CYP2C19, CYP2D6, CYP2E1, CYP3A4, and CYP3A5 (Crespi et al., *The use of heterologously expressed drug metabolizing enzymes-state of the art and prospects for the future*, *Pharm Ther* 84:121-131, 1999).

#### [0593] X.A.3.c. Miscellaneous Membrane Proteins

[0594] In addition to the preceding non-limiting examples, the invention can be applied to the membrane proteins described in U.S. Pat. Nos. 6,335,018 (High molecular weight major outer membrane protein of *Moraxella*); U.S. Pat. No. 6,264,954 (Haemophilus outer membrane protein); U.S. Pat. No. 6,197,543 (Human vesicle membrane protein-like proteins); U.S. Pat. No. 6,121,427 (Major outer membrane protein CD of *Brachyella*); U.S. Pat. Nos. 6,083,743 and 6,013,514 (Haemophilus outer membrane protein); U.S. Pat. No. 6,004,562 (Outer membrane protein B1 of *Moraxella catarrhalis*); U.S. Pat. No. 5,863,764 (DNA encoding a human membrane protein); U.S. Pat. No. 5,861,283 (DNA encoding a limbic system-associated membrane protein); U.S. Pat. No. 5,824,321 (Cloned leptospira outer membrane protein); U.S. Pat. No. 5,821,085 (Nucleotide sequences of a *T. pallidum* rare outer membrane protein); U.S. Pat. No. 5,821,055 (Chlamydia major outer membrane protein); U.S. Pat. No. 5,808,024 (Nucleic acids encoding high molecular weight major outer membrane protein of *Moraxella*); U.S. Pat. No. 5,770,714 (Chlamydia major outer membrane protein); U.S. Pat. No. 5,763,589 (Human mem-

brane protein); U.S. Pat. No. 5,753,459 (Nucleotide sequences of *T. pallidum* rare outer membrane protein); U.S. Pat. No. 5,607,920 (Concanavalin a binding proteins and a 76 kD chondrocyte membrane protein (CMP) from chondrocytes and methods for obtaining same); and U.S. Pat. No. 5,503,992 (DNA encoding the 15 kD outer membrane protein of *Haemophilus influenzae*).

**[0595]** X.B. Membrane Anchoring Domains

**[0596]** A membrane-anchoring domain can be incorporated into a fusion protein of the invention. Non-limiting examples of membrane anchoring domains include those derived from Prostaglandin H2 synthases (PGHS-1 and -2) (Nina et al., Anchoring of a monotopic membrane protein: the binding of prostaglandin H2 synthase-1 to the surface of a phospholipid bilayer, *Eur. Biophys. J.* 29:439-54, 2000; Otto and Smith, Photolabeling of prostaglandin endoperoxide H synthase-1 with 3-trifluoro-3-(m-[<sup>125</sup>I]iodophenyl) diazirine as a probe of membrane association and the cyclooxygenase active site, *J Biol Chem* 271:9906-10, 1996; and Otto and Smith, The orientation of prostaglandin endoperoxide synthases-1 and -2 in the endoplasmic reticulum, *J Biol Chem* 269:19868-75, 1994; those derived from carboxypeptidase E (EC 3.4.17.10) (Fricker et al., Identification of the pH-dependent membrane anchor of carboxypeptidase E (EC 3.4.17.10), *J. Biol. Chem.*, 265, 2476-2482, 1990); and peptide convertase 3 (PC3) (Smeekens et al., Identification of a cDNA encoding a second putative prohormone convertase related to PC2 in AtT20 cells and islets of Langerhans, *Proc Natl Acad Sci USA* 88, 340-344, 1990).

**[0597]** X.C. Transmembrane Domains

**[0598]** A variety of types and examples of transmembrane domain are known. Proteins with up to 12 transmembrane domains are known (Fujiwara et al., Identification of thyroid hormone transporters in humans: different molecules are involved in a tissue-specific manner, *Endocrinology* 2001 142:2005-12; Sharina et al., Mutational analysis of the functional role of conserved arginine and lysine residues in transmembrane domains of the murine reduced folate carrier, *Mol Pharmacol* 2001 59:1022-8). However, the invention is not limited to any particular number of transmembrane domains.

**[0599]** Monotropic ("single pass") domains, which traverse a membrane once, include by way of non-limiting example, those found in receptors for epidermal growth factor (EGF), receptors for tumor necrosis factor (TNF) and the like. Polytopic ("multipass") proteins traverse a membrane two or more times. Non-limiting examples of polytopic proteins are as follows.

**[0600]** Biotropic ("2 passes") membrane proteins include, but are not limited to: EnvZ of *E. coli*; the peroxisomal membrane protein Pex11-1p (Anton et al., ARF- and coatmer-mediated peroxisomal vesiculation, *Cell Biochem Biophys* 2000;32 Spring:27-36); pleiotropic drug ABC transporters of *S. cerevisiae* (Rogers et al., The pleiotropic drug ABC transporters from *Saccharomyces cerevisiae*, *J Mol Microbiol Biotechnol* 2001 3:207-14); and human and rat urate transporters hUAT and rUAT (Lipkowitz et al., Functional reconstitution, membrane targeting, genomic structure, and chromosomal localization of a human urate transporter, *J Clin Invest* 2001 107:1103-15).

**[0601]** Tritropic ("3 pass") membrane proteins include, but are not limited to: the ethylene receptor ETR1 of

Arabidopsis; the Cauliflower Card Expression protein CC 1 (Palmer et al., *A Brassica oleracea* Gene Expressed in a Variety-Specific Manner May Encode a Novel Plant Transmembrane Receptor, *Plant Cell Physiol* 2001 42:404-413); and a splice variant of the mitochondrial membrane protein hMRS3/4 (Li et al., Characterization of a novel human putative mitochondrial transporter homologous to the yeast mitochondrial RNA splicing proteins 3 and 4, *FEBS Lett* 2001 494:79-84).

**[0602]** Tetraspanins or tetraspans are non-limiting examples of membrane proteins with four transmembrane domains. (Levy et al., *J. Biol. Chem.*, 226:14597-14602, 1991; Tomlinson et al., *J. 1 mmol.* 23:136-40, 1993; and Barclay et al., (In) *The Leucocyte antigen factbooks*, Academic press, London, 1993). These proteins are collectively known as the transmembrane 4 superfamily (TM4) because they span the plasma membrane four times. The proteins known to belong to this family include, but are not limited to: mammalian antigen CD9 (MIC3), a protein involved in platelet activation and aggregation; mammalian leukocyte antigen CD37, expressed on B lymphocytes; mammalian leukocyte antigen CD53 (OX-44), which may be involved in growth regulation in hematopoietic cells; mammalian lysosomal membrane protein CD63 (Melanoma-associated antigen ME491; antigen AD1); mammalian antigen CD81 (cell surface protein TAPA-1), which may play an important role in the regulation of lymphoma cell growth; mammalian antigen CD82 (Protein R2; Antigen C33; Kangai 1 (KAI1)), which associates with CD4 or CD8 and delivers costimulatory signals for the TCR/CD3 pathway; mammalian antigen CD151 (SFA-1); Platelet-endothelial tetraspan antigen 3 (PETA-3); mammalian TM4SF2 (Cell surface glycoprotein A15; TALLA-1; MXS1); mammalian TM4SF3 (Tumor-associated antigen CO-029); mammalian TM4SF6 (Tspan-6; TM4-D); mammalian TM4SF7 (Novel antigen 2 (NAG-2); Tspan-4); mammalian Tspan-2; Mammalian Tspan-3 (TM4-A); mammalian Tetraspan NET-5; and *Schistosoma mansoni* and japonicum 23 Kd surface antigen (SM23/SJ23).

**[0603]** Non-limiting examples of membrane proteins with six transmembrane domains include the EBV integral membrane protein LMP-1, and a splice variant of the mitochondrial protein hMRS3/4 (Li et al., Characterization of a novel human putative mitochondrial transporter homologous to the yeast mitochondrial RNA splicing proteins 3 and 4, *FEBS Lett Apr.* 6, 2001;494(1-2):79-84). Proteins with six transmembrane domains also include STEAP (six transmembrane epithelial antigens of the prostate) proteins (Afar et al., U.S. Pat. No. 6,329,503). The prototype member of the STEAP family, STEAP-1, appears to be a type IIIa membrane protein expressed predominantly in prostate cells in normal human tissues. Structurally, STEAP-1 is a 339 amino acid protein characterized by a molecular topology of six transmembrane domains and intracellular N- and C-termini, suggesting that it folds in a "serpentine" manner into three extracellular and two intracellular loops.

**[0604]** Literally hundreds of 7-pass membrane proteins are known. G-protein coupled receptors (GPCRs), including without limitation beta-adreno receptors, adrenergic receptors, EDG receptors, adenosine receptors, B receptors for kinins, angiotensin receptors, and opioid receptors are of particular interest. GPCRs are described in more detail elsewhere herein.

- [0605] A non-limiting example of a protein with 9 transmembrane domains is Lipocalin-1 interacting membrane receptor (Wojnar et al., Molecular cloning of a novel Lipocalin-1 interacting human cell membrane receptor (LIMR) using phage-display, *J Biol Chem* 2001 3; [epub ahead of print]).
- [0606] Proteins with both transmembrane and anchoring domains are known. For example, AMPA receptor subunits have transmembrane domains and one membrane-anchoring domain.
- [0607] A variety of databases that describe known, and software programs that predict, membrane anchoring and transmembrane domains are available to those skilled in the art. See, for example Gcrdb.dba GCRDB [G Protein Coupled Receptor database], Tmbase.dba Tmbase [database of transmembrane domains], Prodom.srv ProDom [Protein domains], Tmap.srv TMAP [Protein transmembrane segments prediction], Tm7.srv TM7 [Retrieval of data on G protein-coupled receptors], and Memsat.sof MEMSAT [transmembrane structure prediction program].
- [0608] Quentin and Fichant (*J Mol Microbiol Biotechnol* 2000 2:501-4, ABCdb: an ABC transporter database) have described a database devoted to the ATP-binding cassette (ABC) protein domains (ABCdb), the majority of which energize the transport of compounds across membranes. In bacteria, ABC transporters are involved in the uptake of a wide range of molecules and in mechanisms of virulence and antibiotic resistance. In eukaryotes, most ABC transporters are involved in drug resistance, and many are associated with diseases. ABCdb can be accessed via the World Wide Web (<http://ir21cb.cnrs-mrs.fr/ABCdb/>). See also Sanchez-Fernandez et al., The *Arabidopsis thaliana* ABC protein superfamily: a complete inventory, *J Biol Chem* May 9, 2001; [epub ahead of print], and Rogers et al., The pleiotropic drug ABC transporters from *Saccharomyces cerevisiae*, *J Mol Microbiol Biotechnol* 2001 April;3(2):207-14.
- [0609] X.D. Functions and Activities of Membrane Proteins
- [0610] Non-limiting examples of membrane proteins include membrane-associated enzymes. Membrane-associated enzymes include but not limited to certain enzymes of the electron transport chain (ETC), antigenic proteins such as the major histocompatibility (MHC) antigens, transport proteins, channels, hormone receptors, cytokine receptors, glucose permeases, gap junction proteins and bacteriorhodopsins.
- [0611] A "transport protein" or "transporter" is a type of membrane protein that allows substances to cross plasma membranes at a rate that is faster than what is found by diffusion alone. Some transport proteins expend energy to move substances (active transport). Many active transport proteins are ATPases (e.g., the Na<sup>+</sup>-K<sup>+</sup> ATPase), or at least bind ATP by virtue of comprising an ATP-binding cassette (ABC) (see, e.g., Rogers et al., The pleiotropic drug ABC transporters from *Saccharomyces cerevisiae*, *J Mol Microbiol Biotechnol* 3:207-14, 2001). Nucleobase transporters are reviewed by De Koning and Diallinas (*Nucleobase Transporters*, *Mol Membr Biol* 17:75-94, 2000).
- [0612] A "channel protein" is a protein that facilitates the diffusion of molecules/ions across lipid membranes by forming a hydrophilic pore or "channel" that provides molecules/ions access through lipid membranes, which are generally hydrophobic. Channels are often multimeric, with the pore being formed by subunit-subunit interactions.
- [0613] A "receptor" is a molecular entity, typically a protein, that is displayed on the surface of a cell. A receptor is characterized by high affinity, often a specific binding of a specific substance, typically resulting in a specific biochemical or physiological effect.
- [0614] A "hormone" is a naturally occurring substance secreted by specialized cells that affects the metabolism or behavior of other cells having receptors for the hormone. Non-limiting examples of hormones having receptors include but are not limited to insulin, cytokines, steroid hormones, histamines, glucagon, angiotensin, catecholamines, low density lipids (LDLs), tumor necrosis factor alpha, tumor necrosis factor beta, estrogen, and testosterone.
- [0615] X.E. G-Protein-Coupled Receptors
- [0616] G protein-coupled receptors (GPCRs) constitute the most prominent family of validated drug targets within biomedical research and are thought to be involved in such diseases and disorders as heart disease, hypertension, cancer, obesity, and depression and other mental illnesses. Over half of approved drugs elicit their therapeutic effects by selectively addressing members of this target family and more than 1000 sequences of the human genome encode for GPCRs containing the classical 7-pass membrane structure characteristic of this family of proteins (Marinissen, M. and J. S. Gutkind, G-protein-coupled receptors and signaling networks: emerging paradigms (Review), *Trends Pharmacol. Sci.* 22: 368-376, 2001). Many pharmacological drug companies are interested in the study of G-coupled proteins. It is possible to co-express a G-coupled protein receptor and its associated G-protein to study their pharmacological characteristics (Strosberg and Marullo, Functional expression of receptors in microorganisms. *TIPS*, 1992. 13: 95-98).
- [0617] G-protein-coupled receptors (GPCRs) are reviewed by Marinissen, M. and J. S. Gutkind, G-protein-coupled receptors and signaling networks: emerging paradigms. *Trends Pharmacol. Sci.* 22: 368-376, 2001; Sautel and Milligan, Molecular manipulation of G-protein-coupled receptors: a new avenue into drug discovery, *Curr Med Chem* 2000 889-96; Hibert et al., This is not a G protein-coupled receptor, *Trends Pharmacol Sci* 1993, 14:7-12; Wilson et al., Orphan G-protein-coupled receptors: the next generation of drug targets?, *Br J Pharmacol* 1998, 125: 1387-92; Roth et al., G protein-coupled receptor (GPCR) trafficking in the central nervous system: relevance for drugs of abuse, *Drug Alcohol Depend* 1998, 51:73-85; Ferguson and Caron, G protein-coupled receptor adaptation mechanisms, *Semin Cell Dev Biol* 1998, 9:119-27; Wank, G protein-coupled receptors in gastrointestinal physiology. I. CCK receptors: an exemplary family, *Am J Physiol* 1998, 274:G607-13; Rohrer and Kobilka, G protein-coupled receptors: functional and mechanistic insights through altered gene expression. (Review), *Physiol Rev* 1998, 78:35-52; and Larhammar et al., The receptor revolution—multiplicity of G-protein-coupled receptors. (Review), *Drug Des Discov* 1993, 9:179-88.
- [0618] GPCR localization and regulation has been studied using GFP-comprising fusion proteins (Kallal and Benovic,

Using green fluorescent proteins to study G-protein-coupled receptor localization and trafficking. (Review), *Trends Pharmacol Sci* 2000 21:175-80; and Ferguson, Using green fluorescent protein to understand the mechanisms of G-protein-coupled receptor regulation. (Review), *Braz J Med Biol Res* 1998, 31:1471-7; and by using chimeric GPCRs (Milligan and Rees, Chimeric G alpha proteins: their potential use in drug discovery. (Review), *Erratum in: Trends Pharmacol Sci* 1999 June; 20(6):252.

[0619] GPCRs belong to a superfamily of at least 6 families of receptors, the most important of which is the main family, A. Members of the membrane protein gene superfamily of GPCRs have been characterized as having seven putative transmembrane domains. The transmembrane domains are believed to represent transmembrane alpha-helices connected by extracellular or cytoplasmic loops. A functional G-protein is a trimer which consists of a variable alpha subunit coupled to much more tightly-associated and constant beta and gamma subunits, although G-protein independent actions have been postulated (Marinissen, M. and J. S. Gutkind, G-protein-coupled receptors and signaling networks: emerging paradigms. *Trends. Pharmacol. Sci.* 22: 368-376, 2001 Review). A variety of ligands have been identified which function through GPCRs. In general, binding of an appropriate ligand (e.g., bioactive lipids, ions, bioactive amines, photons, odorants, hormones, neurotransmitters, peptides, nucleosides, etc.) to a GPCR leads to the activation of the receptor. G-protein coupled receptors include a wide range of biologically active receptors, such as hormone, viral, growth factor and neuroreceptors. Typically, activation of a GPCR initiates the regulatory cycle of a corresponding G-protein. This cycle consists of GTP exchange for GDP, dissociation of the alpha and beta/gamma subunits, activation of the second messenger pathway by a complex of GTP and the alpha subunit of the G-protein, and return to the resting state by GTP hydrolysis via the innate GTPase activity of the G-protein alpha subunit A.

[0620] GPCRs include, without limitation, dopamine receptors which bind to neuroleptic drugs used for treating psychotic and neurological disorders. Other examples of members of this family include calcitonin, adrenergic, endothelin, cAMP, adenosine, muscarinic, acetylcholine, serotonin, histamine, thrombin, kinin, follicle stimulating hormone, opsins and rhodopsins, odorant, cytomegalovirus receptors, and the like.

[0621] Most GPCRs have single conserved cysteine residues in each of the first two extracellular loops which form disulfide bonds that are believed to stabilize functional protein structure. The seven transmembrane regions, each comprising conserved hydrophobic stretches of about 20 to 30 amino acids, are designated as TM1, TM2, TM3, TM4, TM5, TM6, and TM7. TM3 is also implicated in signal transduction.

[0622] Although not wishing to be bound by any particular theory, it is believed that GPCRs participate in cell signaling through their interactions with heterotrimeric G-proteins composed of alpha, beta and gamma subunits (Marinissen, M. and J. S. Gutkind, G-protein-coupled receptors and signaling networks: emerging paradigms. *Trends. Pharmacol. Sci.* 22:368-376, 2001). In some aspects of the invention, GPCRs and homologs are displayed on the surfaces of micelles.

[0623] X.F. EDG Receptors and Other Sphingolipid-Binding Receptors

[0624] The Endothelial Differentiation Gene (EDG) receptor family includes but is not limited to eight presently known GPCRs that have a high affinity to lipid ligands (Lynch et al., *Life on the edge. Trends Pharmacol. Sci.*, 1999, 20: 273-5). These transmembrane receptors are found in several different tissues in different species. EDG receptors have been shown to be involved in calcium mobilization, activation of mitogen-activated protein kinase, inhibition of adenylate cyclase activation, and alterations of the cytoskeleton. The EDG family is divided into two different groups based on homology and ligand specificity. The EDG 2, 4, and 7 receptors are specific for the ligand lysophosphatidic acid (LPA) (An et al., *Signaling Mechanism and molecular characteristics of G protein-coupled receptors for lysophosphatidic acid and sphingosine 1-phosphate. J. Cell Biochem*, 30/31:147-157, 1998; Goetzl et al., *Distinctive expression and functions of the type 4 endothelial differentiation gene-encoded G protein-coupled receptor for lysophosphatidic acid in ovarian cancer. Cancer Res.*, 59:5370-5, 1999). In contrast, EDG 1, 3, and 5 bind sphingosine-1-phosphate (S1P) (Zhang et al., *Comparative analysis of three murine G-protein coupled receptors activated by sphingosine-1-phosphate. Gene*, 227:89-99, 1999). EDG-6 is believed to interact with S1P (Yamazaki et al., *Edg-6 as a putative sphingosine 1-phosphate receptor coupling to Ca<sup>2+</sup> signaling pathway. Biochem Phys Res Com*, 268:583-589, 2000).

[0625] Receptors that bind S1P and other sphingolipids are used in one aspect of the invention (for a review of some S1P-binding receptors, see Spiegel et al., *Biophys. Acta* 1484:107-116, 2000). Such receptors include but are not limited to members of the EDG family of receptors (a.k.a. 1pA receptors, Chun, *Crit. Rev. Neuro.* 13:151-168, 1999), and isoforms and homologs thereof such as NRG1 and AGR16.

[0626] EDG-1 was the first identified member of a class of G protein-coupled endothelial-derived receptors (EDG). Non-limiting examples of other EDG family members that also bind S1P include EDG-3 (a.k.a. ARG 16; the rat homolog of EDG-3 is designated H218), EDG-5, EDG-6 and EDG-8. **For reviews, see Goetzl et al., *Adv. Exp. Med. Biol.* 469:259-264, 1999; and Chun et al., *Cell. Biochem. Biophys.* 30:213-242, 1999).**

[0627] EDG-1 is described by Lee et al., (*Ann. NY Acad. Sci.* 845:19-31, 1998). Liu and Hla, The mouse gene for the inducible G-Protein-coupled receptor edg-1. *Genomics*, 1997, 43: p.15-24. Human EDG-1c genes and proteins are described in published PCT application WO 99/46277 to Bergsma et al.

[0628] EDG-3 is described by Okamoto et al. (*Biochem. Biophys. Res. Commun.* 260:203-208, 1999) and An et al. (*FEBS Letts.* 417:279-282, 1997). See also An et al., *J. Biol. Chem.* 275:288-296, 2000.

[0629] EDG-5 human and mammalian genes are described in U.S. Pat. No. 6,057,126 to Munroe et al. and published PCT application WO 99/33972 to Munroe et al. The rat homolog, H218, is described in U.S. Pat. No. 5,585,476 to MacLennan et al. Van Brocklyn et al., *J. Biol. Chem.* 274:4626-4632, 1999; and Gonda et al., *Biochem. J.* 337:67-75, 1999. See also An et al., *J. Biol. Chem.* 275:288-296, 2000.

[0630] EDG-6 is described by Graler et al. (*Genomics* 53:164-169, 1998), Yamazaki et al. (*Biochem. Biophys. Res. Commun.* 268:583-589, 2000), and Van Brocklyn et al. (Sphingosine-1-phosphate is a ligand for the G protein-coupled receptor EDG-6, *Blood* 95:2624-9, 2000).

[0631] EDG-8 from rat brain is described by Im et al., (*J. Biol. Chem.* 275:14281-14286, 2000). Homologs of EDG-8 from other species, including humans, may also be used in the present invention.

[0632] The Mil receptor (Mil is an abbreviation for "miles apart") binds S1P and regulates cell migration during vertebrate heart development. The Mil receptor of Zebrafish is described by Mohler et al. (*J. Immunol.* 151:1548-1561, 1993). Another S1P receptor is NRG1 (nerve growth factor regulated gene-1), the rat version of which has been identified (Glickman et al., *Mol. Cel. Neurosci.* 14:141-152, 1999).

[0633] Receptors that bind sphingosylphosphoryl choline (SPC) are also used in this aspect of the invention. Such receptors include but are not limited to members of the SCaMPER family of receptors (Mao et al., *Proc. Natl. Acad. Sci. U.S.A.* 93:1993-1996, 1996; Betto et al., *Biochem. J.* 322:327-333, 1997). Some evidence suggests that EDG-3 may bind SPC in addition to S1P (Okamoto et al., *Biochem. Biophys. Res. Commun.* 260:203-208, 1999). Derivatives of EDG-3 that bind both S1P and SPC are used in one aspect of the invention.

[0634] Receptors that bind lysophosphatidic acid may be used in the present invention. These include EDG-2 (LPA1), EDG-4 (LPA2), EDG-7 (LPA3). See Moller et al., Expression and function of lysophosphatidic acid receptors in cultured rodent microglial cells, *J Biol Chem* May 4, 2001 [epub ahead of print]; Fukushima and Chun, The LPA receptors, Prostaglandins 64(1-4):21-32, 2001; Contos and Chun, The mouse lp(A3)/Edg7 lysophosphatidic acid receptor gene: genomic structure, chromosomal localization, and expression pattern, *Gene* 267:243-53, 2001; Schulte et al., Lysophosphatidic acid, a novel lipid growth factor for human thyroid cells: over-expression of the high-affinity receptor edg4 in differentiated thyroid cancer, *Int J Cancer* 92249-56, 2001; Kimura et al., Two novel Xenopus homologs of mammalian LP(A1)/EDG-2 function as lysophosphatidic acid receptors in Xenopus oocytes and mammalian cells, *J Biol Chem* 276:15208-15, 2001; and Swarthout and Walling, Lysophosphatidic acid: receptors, signaling and survival (Review), *Cell Mol Life Sci* 57:1978-85, 2000.

[0635] Examples of lysophospholipid receptors including, but not limited to EDG proteins, are disclosed in Fukushima et al. (Lysophospholipid receptors. *Annu. Rev. Pharmacol. Toxicol.* 41:507-534, 2001) Malek and Lee (Nrg-1 Belongs to the Endothelial Differentiation Gene Family of G Protein-coupled Sphingosine-1-phosphate Receptors, *J. Biol. Chem.* 276:5692-5699, 2001), Hla et al. (Sphingosine-1-phosphate signaling via the EDG-1 family of G-protein-coupled receptors (Review), *Ann NY Acad Sci* 905:16-24, 2000; Chun, Lysophospholipid receptors: implications for neural signaling (Review), *Crit Rev Neurobiol* 13:151-68, 1999); and Chun et al. (A growing family of receptor genes for lysophosphatidic acid (LPA) and other lysophospholipids (LPs) (Review), *Cell Biochem Biophys* 30:213-42, 1999).

[0636] XI. Recombinant DNA Expression

[0637] In order to achieve recombinant expression of a fusion protein, an expression cassette or construct capable of

expressing a chimeric reading frame is introduced into an appropriate host cell to generate an expression system. The expression cassettes and constructs of the invention may be introduced into a recipient eubacterial or eukaryotic cell either as a nonreplicating DNA or RNA molecule, which may be a linear molecule or, more preferably, a closed covalent circular molecule. Since such molecules are incapable of autonomous replication, the expression of the gene may occur through the transient expression of the introduced sequence. Alternatively, permanent expression may occur through the integration of the introduced DNA sequence into the host chromosome.

[0638] XI.A. Recombinant DNA Expression Systems

[0639] A variety of eubacterial recombinant DNA expression systems may be used to produce the fusion proteins of the invention. Host cells that may be used in the expression systems of the present invention are not strictly limited, provided that they are suitable for use in the expression of the fusion protein of interest and can produce minicells. Non-limiting examples of recognized eubacterial hosts that may be used in the present invention include bacteria such as *E. coli*, *Bacillus*, *Streptomyces*, *Pseudomonas*, *Salmonella*, *Serratia*, and the like.

[0640] Eubacterial expression systems utilize plasmid and viral (bacteriophage) expression vectors that contain replication sites and control sequences derived from a species compatible with the host may be used. Suitable phage or bacteriophage vectors include  $\lambda$ gt10,  $\lambda$ gt11 and the like. Suitable virus vectors may include pMAM-neo, pKRC and the like. Appropriate eubacterial plasmid vectors include those capable of replication in *E. coli* (such as, by way of non-limiting example, pBR322, pUC 118, pUC 119, ColE1, pSCO101, pACYC 184,  $\pi$ VX. See "Molecular Cloning: A Laboratory Manual" 1989). *Bacillus* plasmids include pC194, pC221, pT127, and the like (Gryczan, In: *The Molecular Biology of the Bacilli*, Academic Press, NY, pp. 307-329, 1982). Suitable *Streptomyces* plasmids include p1J101 (Kendall et al., *J. Bacteriol.* 169:4177-4183, 1987), and *Streptomyces* bacteriophages such as C31 (Chater et al., In: *Sixth International Symposium on Actinomycetales Biology*, Akademiai Kiado, Budapest, Hungary, pp. 45-54, 1986). *Pseudomonas* plasmids are reviewed by John et al. (*Rev. Infect. Dis.* 8:693-704, 1986), and Izaki (*Jpn. J. Bacteriol.* 33:729-742, 1978). See also Brent et al., "Vectors Derived From Plasmids," Section II, and Lech et al. "Vectors derived from Lambda and Related Bacteriophages" Section III, in Chapter 1 of *Short Protocols in Molecular Biology*, 2nd Ed., Ausubel et al., eds., John Wiley and Sons, New York, 1992, pages 1-13 to 1-27; Lech et al. "Vectors derived from Lambda and Related Bacteriophages" Section III and Id. pages 1-28 to page 1-52.

[0641] To express a protein, including but not limited to a fusion protein, in a eubacterial cell, it is necessary to operably link the ORF encoding the protein to a functional eubacterial or viral promoter. Such promoters may be either constitutive or, more preferably, regulatable (i.e., inducible or derepressible). Examples of constitutive promoters include the int promoter of bacteriophage lambda, the bla promoter of the beta-lactamase gene sequence of pBR322, and the cat promoter of the chloramphenicol acetyl trans-

ferase gene sequence of pPR325, and the like. Examples of inducible eubacterial promoters include the major right and left promoters of bacteriophage lambda (PL and PR), the *trp*, *recA*, *lacZ*, *lacI*, and *gal* promoters of *E. coli*, the alpha-amylase (Ulmanen et al., *J. Bacteriol.* 162:176-182, 1985) and the sigma-28-specific promoters of *B. subtilis* (Gilman et al., *Gene Sequence* 32:11-20, 1984), the promoters of the bacteriophages of *Bacillus* (Gryczan, in: *The Molecular Biology of the Bacilli*, Academic Press, Inc., NY, 1982), and *Streptomyces* promoters (Ward et al., *Mol. Gen. Genet.* 203:468-478, 1986). Eubacterial promoters are reviewed by Glick (*Ind. Microbiol.* 1:277-282, 1987), Cenatiempo (*Biochimie* 68:505-516, 1986), and Gottesman (*Ann. Rev. Genet.* 18:415-442, 1984).

[0642] Proper expression also requires the presence of a ribosome-binding site upstream of the gene sequence-encoding sequence. Such ribosome-binding sites are disclosed, for example, by Gold et al. (*Ann. Rev. Microbiol.* 35:365-404, 1981). The selection of control sequences, expression vectors, transformation methods, and the like, are dependent on the type of host cell used to express the gene. As used herein, "cell", "cell line", and "cell culture" may be used interchangeably and all such designations include progeny. Thus, the words "transformants" or "transformed cells" include the primary subject cell and cultures derived therefrom, without regard to the number of transfers. It is also understood that all progeny may not be precisely identical in DNA content, due to deliberate or inadvertent mutations. However, as defined, mutant progeny have the same functionality as that of the originally transformed cell.

[0643] Mammalian expression systems utilize host cells such as HeLa cells, cells of fibroblast origin such as VERO or CHO-K1, or cells of lymphoid origin and their derivatives. Preferred mammalian host cells include SP2/0 and J558L, as well as neuroblastoma cell lines such as IMR 332, which may provide better capacities for correct post-translational processing. Non-limiting examples of mammalian extrachromosomal expression vectors include pCR3.1 and pDNA3.1, and derivatives thereof including but not limited to those that are described by and are commercially available from Invitrogen (Carlsbad, Calif.).

[0644] Several expression vectors are available for the expression of polypeptides in mammalian host cells. A wide variety of transcriptional and translational regulatory sequences may be employed, depending upon the nature of the host. The transcriptional and translational regulatory signals may be derived from viral sources, such as adenovirus, bovine papilloma virus, cytomegalovirus (CMV), simian virus, or the like, where the regulatory signals are associated with a particular gene sequence which has a high level of expression. Alternatively, promoters from mammalian expression products, such as actin, collagen, myosin, and the like, may be employed. Transcriptional initiation regulatory signals may be selected which allow for repression or activation, so that expression of the gene sequences can be modulated. Of interest are regulatory signals that are temperature-sensitive since, by varying the temperature, expression can be repressed or initiated, or are subject to chemical (such as metabolite) regulation.

[0645] Preferred eukaryotic plasmids include, for example, BPV, vaccinia, SV40, 2-micron circle, and the like, or their derivatives. Such plasmids are well known in the art

(Botstein et al., *Miami Wntr. Symp.* 19:265-274, 1982; Broach, in: *The Molecular Biology of the Yeast Saccharomyces: Life Cycle and Inheritance*, Cold Spring Harbor Laboratory, Cold Spring Harbor, N.Y., p. 445-470, 1981; Broach, *Cell* 28:203-204, 1982; Bollon et al., *J. Clin. Hematol. Oncol.* 10:39-48, 1980; Maniatis, In: *Cell Biology: A Comprehensive Treatise*, Vol. 3, *Gene Sequence Expression*, Academic Press, NY, pp. 563-608, 1980).

[0646] Expression of polypeptides in eukaryotic hosts generally involves the use of eukaryotic regulatory regions. Such regions will, in general, include a promoter region sufficient to direct the initiation of RNA synthesis. Preferred eukaryotic promoters include, for example, the promoter of the mouse metallothionein I gene sequence (Hamer et al., *J. Mol. Appl. Gen.* 1:273-288, 1982); the TK promoter of Herpes virus (McKnight, *Cell* 31:355-365, 1982); the SV40 early promoter (Benoist et al., *Nature (London)* 290:304-31, 1981); and the yeast *gal4* gene sequence promoter (Johnston et al., *Proc. Natl. Acad. Sci. (USA)* 79:6971-6975, 1982; Silver et al., *Proc. Natl. Acad. Sci. (USA)* 81:5951-5955, 1984).

[0647] Expression sequences and elements are also required for efficient expression. Non-limiting examples include Kozak and IRES elements in eukaryotes, and Shine-Delgarno sequences in prokaryotes, which direct the initiation of translation (Kozak, *Initiation of translation in prokaryotes and eukaryotes*. *Gene*, 1999. 234: 187-208; Martinez-Salas et al., *Functional interactions in internal translation initiation directed by viral and cellular IRES elements*, *Jour. of Gen. Virol.* 82:973-984, 2001); enhancer sequences; optioanal sites for repressor and inducers to bind; and recognition sites for enzymes that cleave DNA or RNA in a site-specific manner. Translation of mRNA is generally initiated at the codon which encodes the first methionine; if so, it is preferable to ensure that the linkage between a eukaryotic promoter and a preselected ORF does not contain any intervening codons that encode a methionine (i.e., AUG). The presence of such codons results either in the formation of a fusion protein with an uncharacterized N-terminal extension (if the AUG codon is in the same reading frame as the ORF) or a frame-shift mutation (if the AUG codon is not in the same reading frame as the ORF).

[0648] XI.B. Expression of Membrane Proteins Presently, the most commonly used expression systems for the expression of integral membrane proteins are eukaryotic and eubacterial whole cell expression systems. Although minicells have been used to express several eubacterial membrane proteins, the production of non-eubacterial membrane proteins has not been reported. One aspect of the invention is the discovery that the minicell expression system can be made to express and preferably display integral membrane proteins from non-eubacterial organisms.

[0649] Some commonly used expression systems include in vitro systems, such as the Rabbit Reticulocyte Lysate System and *E. coli* S30 Extract System (both available from Promega) (Zubay, *Methods Enz.* 65:856, 1980) and in vivo systems, such as eukaryotic cell culture expression, and bacterial expression systems. Although this is not an exhaustive list, these systems are representative.

[0650] The Rabbit Reticulocyte Lysate system utilizes a cell lysate that contains all the enzymes required for transcription and translation to drive protein expression, and is



a good in vitro system for producing small amounts of labeled and unlabeled protein. However, this system is not well-suited for the production of large quantities of proteins and is limited to soluble proteins as there are no membranes in which to incorporate membrane proteins.

[0651] In eukaryotic cell culture systems, expression vectors suited for expression in host eukaryotic cells are transfected into cultured cells and protein is translated from mRNA produced from the vector DNA template Kaufman, Overview of vector design for mammalian gene expression. Mol Biotechnol, 2001. 16: 151-160; Lee, et al., Heterologous gene expression in avian cells: Potential as a producer of recombinant proteins. J Biomed Sci, 1999. 6: 8-17; Voorma et al., Initiation of protein synthesis in eukaryotes. Mol Biol Rep, 1994. 19: 139-45). Cells can then either be harvested to prepare at least partially purified proteins or proteins produced from the expression element can be studied in the host cell environment.

[0652] Regarding membrane proteins, such systems have limitations. Primary cell lines are difficult to maintain and are short lived. Immortalized cell lines divide indefinitely, but have been altered in many ways and can be unpredictable. The transfection efficiency is very low in most eukaryotic cells and some cell types are refractory to transformation. Moreover, other proteins are expressed in these cells along with the protein of interest. This can cause difficulties when performing certain experiments and when attempting to immunoprecipitate the protein. Good experimental data are difficult to obtain from studies such as binding assays (because of high background due to endogenous proteins), and crystal determination of protein structure (because it is difficult to obtain enough purified protein to efficiently form crystals).

[0653] Bacterial expression systems are generally similar to that of the eukaryotic expression systems in that they both use the host cell enzymes to drive protein expression from recombinant expression vectors (Cornelis, P., Expressing genes in different *Escherichia coli* compartments. Curr Opin Biotechnol, 2000. 11: p. 450-454; Laage and Langosch, Strategies for prokaryotic expression of eukaryotic membrane proteins. Traffic, 2001. 2: 99-104; Pines, O. and M. Inouye, Expression and secretion in *E. coli*. Mol Biotechnol, 1999. 12: 25-34).

[0654] In bacterial expression systems, bacterial cells are transformed with expression elements, and transcription and translation is driven from a bacterial promoter. Bacteria divide very rapidly and are easy to culture; it is relatively easy to produce a large number of bacteria in a short time. Moreover, incorporation of expression elements vector into bacterial cells is efficient. Transformed cells can be isolated that arise from a single bacterium. Cultures of transformed cells are thus genetically identical and all cells in the culture will contain the expression element. However, there are proteins that are not suitable for expression in bacteria because of differences between eukaryotic cells and bacterial cells in transcription, translation, and post-translational modification.

[0655] The *E. coli* whole cell expression system has been used to express functional integral membrane proteins. For a review, see Strosberg, Functional expression of receptors in microorganisms. TiPS, 1992. 13: 95-98. Examples of mammalian integral membrane proteins that have been

expressed in *Escherichia coli* include rat alpha-2B-adrenoceptors (Xia et al., Functional expression of rat  $\beta$ 2B-adrenoceptor in *E. coli*. Euro J. Pharma, 1993. 246: 129-133) and the human beta2-adrenergic receptor (Marullo et al., Human  $\beta$ 2-adrenergic receptors expressed in *Escherichia coli* membranes retain their pharmacological properties. Proc. Natl. Acad. Sci. USA, 1988. 85: 7551-7555). In some of these studies, the integral membrane proteins were not only expressed in *E. coli* expression systems, but also retained their pharmacological properties. This allows for binding studies to be performed with minimal background signal ("noise") from host cell proteins. It has also been shown that signal sequences (the short hydrophobic amino acid sequence at the N-terminus of integral membrane proteins that signals the transport of the protein to the membrane) from mammalian cells may be functional in the *E. Coli* system.

[0656] As is discussed herein, the expression of membrane proteins such as GPCRs, ion channels, and immuno-receptors in minicells, and their incorporation into the membranes thereof, allows for the study and use of such non-eubacterial membrane proteins. The minicell system of the invention is particularly well-suited for the study and expression of EDG proteins because of the lipid nature of the ligands for these receptors. The identification of ligand binding kinetics and biochemistry of these receptors because of the physiochemical properties of the lipid ligands (LPA and S1P), which results in high non-specific binding (Lee et al., Sphingosine-1-phosphate is a ligand for the G protein-coupled receptor EDG-1. Science, 1998. 279: 1552-1555; Van Brocklyn et al., Sphingosine-1-phosphate is a ligand for the G protein-coupled receptor EDG-6. Blood, 2000. 95: 2624-2629; Liu et al., Edg-1, the G protein-coupled receptor for sphingosine-1-phosphate, is essential for vascular maturation. J. Clin. Investigation, 2000. 106: 951-961).

[0657] It is believed, for example, that in the case of the ion channels, the minicell expression system is less cumbersome than procedures that are presently used to study properties of ion channels, such as, e.g., reconstitution studies (Montal, Molecular anatomy and molecular design of channel proteins. FASEB J., 1990. 4: p. 2623-2635). Ionic conditions both inside and outside of minicells can be manipulated in various ways, and the properties of an ion channel that is expressed in a minicell, and factors that activate or modulate the activities of the channel, can be studied. Binding and kinetic studies are performed on ligand mediated ion channels. This type of study is enhanced when the ion channel is able to interact specifically with its ligand and has a low background of non-specific binding from the endogenous proteins. This can be accomplished by making the minicells into protoplasts or poroplasts in which the ligand-activated ion channels in the inner membrane are exposed to the external environment and have better access to their specific ligand.

[0658] A "recombinant expression system" (or simply "expression system") is one that directs the production of exogenous gene products in a host cell or minicell of choice. By "expressed" it is meant that a gene product of interest (which can be a protein or nucleic acid) is produced in the expression system of choice.

[0659] Host cells (and/or minicells) harboring an expression construct are components of expression systems. An

“expression vector” is an artificial nucleic acid molecule into which an exogenous ORF encoding a protein, or a template of a bioactive nucleic acid can be inserted in such a manner so as to be operably linked to appropriate expression sequences that direct the expression of the exogenous gene. By the term “operably linked” it is meant that the part of a gene that is transcribed is correctly aligned and positioned with respect to expression sequences that promote, are needed for and/or regulate this transcription. The term “gene product” refers to either a nucleic acid (the product of transcription, reverse transcription, or replication) or a polypeptide (the product of translation) that is produced using the non-vector nucleic acid sequences as a template.

[0660] In some applications, it is preferable to use an expression construct that is an episomal element. If the episomal expression construct expresses (or, preferably in some applications, over-expresses) a an ORF that has been incorporated into the episomal expression construct, the minicells will direct the production of the polypeptide encoded by the ORF. At the same time, any mRNA molecules transcribed from a chromosomal gene prior to minicell formation that have been transferred to the minicell are degraded by endogenous RNases without being replaced by new transcription from the (absent) bacterial chromosome.

[0661] Chromosomally-encoded mRNAs will not be produced in minicells and will be “diluted” as increasing amounts of mRNAs transcribed from the episomal element are generated. A similar dilution effect is expected to increase the relative amount of episomally-generated proteins relative to any chromosomally-encoded proteins present in the minicells. It is thus possible to generate minicells that are enriched for proteins encoded by and expressed from episomal expression constructs.

[0662] Although by no means exhaustive, a list of episomal expression vectors that have been expressed in eubacterial minicells is presented in Table 4.

[0663] It is also possible to transform minicells with exogenous DNA after they have been prepared or separated from their parent cells. For example, phage RNA is produced in minicells after infection by lambda phage (Witkiewicz and Taylor, Ribonucleic acid synthesis after adsorption of the bacteriophage lambda on *Escherichia coli* minicells, Acta Microbiol Pol A 7:21-4, 1975), even though replication of lambda phage may not occur in minicells (Witkiewicz and Taylor, The fate of phage lambda DNA in lambda-infected minicells, Biochim Biophys Acta 564:31-6, 1979).

[0664] Because it is the most characterized minicell-producing species, many of these episomal elements have been examined in minicells derived from *E. coli*. It is understood by practitioners of the art, however, that many episomal elements that are expressed in *E. coli* also function in other eubacterial species, and that episomal expression elements for minicell systems in other species are available for use in the invention disclosed herein.

[0665] In one aspect of the invention, eukaryotic and archaeobacterial minicells are used for expression of membrane proteins, particularly in instances where such desirable proteins have enhanced or altered activity after they undergo post-translational modification processes such as phosphorylation, proteolysis, myristylation, GPI anchoring

and glycosylation. Expression elements comprising expression sequence operably linked to ORFs encoding the membrane proteins of interest are transformed into eukaryotic cells according to methods and using expression vectors known in the art. By way of non-limiting example, primary cultures of rat cardiomyocytes have been used to produce exogenous proteins after transfection of expression elements therefor by electroporation (Nakajima et al., Expression and characterization of Edg-1 receptors in rat cardiomyocytes: Calcium deregulation in response to sphingosine-1-phosphate, Eur. J. Biochem. 267: 5679-5686, 2000).

[0666] Yeast cells that produce minicells are transformed with expression elements comprising an ORF encoding a membrane protein operably linked to yeast expression sequences. Cells that harbor a transferred expression element may be selected using a gene that is part of the expression element that confers resistant to an antibiotic, e.g., neomycin.

[0667] Alternatively, in one aspect of the invention, bacterial minicells are prepared that contain expression elements that are prepared from shuttle vectors. A “shuttle vector” has sequences required for its replication and maintenance in cells from two different species of organisms, as well as expression elements, at least one of which is functional in bacterial cells, and at least one of which is functional in yeast cells. For example, *E. coli*-yeast shuttle vectors are known in the art and include, by way of non-limiting example, those derived from Yip, Yrp, Ycp and Yep. Preferred *E. coli*-yeast shuttle vectors are episomal elements that can segregate into yeast minicells (i.e., Yrp, Ycp and Yep. Particularly preferred are expression vectors of the Yep (yeast episomal plasmid) class, and other derivatives of the naturally occurring yeast plasmid known as the 2  $\mu$ m circle. The latter vectors have relatively high transformation frequencies and are stably maintained through mitosis and meiosis in high copy number.

TABLE 4

Episomal Elements That Segregate Into <i>Escherichia coli</i> Minicells	
EPISOMAL ELEMENT	REFERENCES
<u>Plasmids</u>	
R6K, R1DRD19	Nesvera et al., Folia Microbiol. (Praha) 23: 278–285 (1978)
PSC101	Fox et al., Blood 69: 1394–1400 (1987)
PBR322	Fox et al., Blood 69: 1394–1400 (1987)
F element	Cohen et al., Proc. Natl. Acad. Sci. 61: 61–68 (1968); Khachatourians G. G., Biochim. Biophys. Acta. 561: 294–300 (1979)
NR1	Hochmannova et al., Folia Microbiol. (Praha) 26: 270–276
R6 $\delta$ 1	Hochmannova et al., Folia Microbiol. (Praha) 26: 270–276
PTTQ18	Rigg et al., Arch. Oral. Biol. 45: 41–52 (2000)
PGPR2.1	Rigg et al., Arch. Oral. Biol. 45: 41–52 (2000); expresses cell surface antigen of <i>P. gingivalis</i>
“mini-plasmid” derivative of RK2	Firshein et al., J. Bacteriol. 150: 1234–1243 (1982)
ColE1	Rashtchian et al., J. Bacteriol. 165: 82–87 (1986); Witkiewicz et al., Acta. Microbiol. Pol. A 7: 21–24 (1975)
PSC101	Rashtchian et al., J. Bacteriol. 165: 82–87 (1986); Curtiss, Roy, III, U.S. Pat. No. 4,190,495; Issued Feb. 26, 1980

TABLE 4-continued

Episomal Elements That Segregate Into <i>Escherichia coli</i> Minicells	
EPISOMAL ELEMENT	REFERENCES
pACYC184	Chang et al., J. Bacteriol. 134: 1141-1156 (1978); Rose, Nucleic Acids Res 16: 355 (1988)
Co1Ib, Co1Ib7	Skorupska et al., Acta. Microbiol. Pol. A 8: 17-26 (1976)
pUC19	Heighway et al., Nucleic Acids Res. 17: 6893-6901 (1989)
R-plasmid	Hochmannova et al., Folia Microbial. (Praha) 25: 11-15 (1980)
PCR1	Hollenberg et al., Gene 1: 33-47 (1976); yeast shuttle vector
<u>Bacteriophage</u>	
Lambda	Witkiewicz et al., Acta. Microbial. Pal. A 7: 21-24 (1975)
M13	Staudenbauer et al., Mol. Gen. Genet. 138: 203-212 (1975)
T7	Libby, Mech Ageing Dev. 27: 197-206 (1984)
P1	Curtiss, Roy, III, U.S. Pat. No. 4,190,495; Issued Feb. 26, 1980; J Bacteriol 1995; 177: 2381-6, Partition of P1 plasmids in <i>Escherichia coli</i> mukB chromosomal partition mutants, Funnell and Gagnier.

[0668] For expression of membrane proteins, and/or other proteins of interest in the recipient cell, ORFs encoding such proteins are operably linked to eukaryotic expression sequences that are appropriate for the recipient cell. For example, in the case of *E. coli*-yeast shuttle vectors, the ORFs are operably linked to expression sequences that function in yeast cells and/or minicells. In order to assess the effectiveness of a gene delivery vehicle, or a gene therapy expression element, an ORF encoding a detectable polypeptide (e.g., GFP, beta-galactosidase) is used. Because the detectable polypeptide is operably linked to eukaryotic expression elements, it is not expressed unless it has been transferred to its recipient (eukaryotic) cell. The signal from the detectable polypeptide thus correlates with the efficiency of gene transfer by a gene delivery agent, or the degree of expression of a eukaryotic expression element.

[0669] Gyuris and Duda (High-efficiency transformation of *Saccharomyces* cells by bacterial minicell protoplast fusion, Mol Cel Biol 6:329507, 1986) allegedly demonstrated the transfer of plasmid molecular by fusing minicell protoplasts with yeast protoplasts. Gyuris and Duda state that 10% of *Saccharomyces cerevisiae* cells were found to contain transforming DNA sequences. However, the plasmids did not contain eukaryotic expression elements, were not shuttle vectors, and genetic expression of the plasmids in yeast cells was not examined.

[0670] XII. Uses of Minicells in Research

[0671] XII.A. In General

[0672] The minicells of the invention can be used in research applications such as, by way of non-limiting example, proteomics, physiology, chemistry, molecular biology, physics, genetics, immunology, microbiology, proteomics, virology, pathology, botany, and neurobiology. Research applications include but are not limited to protein-ligand binding studies, competitive inhibition studies, structural studies, protein interaction studies, transfection, signaling

studies, viral interaction studies, ELISA, antibody studies, gel electrophoresis, nucleotide acid) applications, peptide production, cell culture applications, cell transport studies, isolation and separation studies, chromatography, labeling studies, synthesis of chemicals, chemical cross linking, flow cytometry, nanotechnology, micro switches, micro-machines, agricultural studies, cell death studies, cell-cell interactions, proliferation studies, and protein-drug interactions. Minicells are applicable to research applications involving, by way of non-limiting example, the elucidation, manipulation, production, replication, structure, modeling, observations, and characterization of proteins.

[0673] The types of proteins that can be involved in research applications of minicells can be either soluble proteins or membrane bound proteins, and include but are not limited to receptors (e.g., GPCRs, sphingolipid receptors, neurotransmitter receptors, sensory receptors, growth factor receptors, hormone receptors, chemokine receptors, cytokine receptors, immunological receptors, and complement receptors, FC receptors), channels (e.g., potassium channels, sodium channels, calcium channels.), pores (e.g., nuclear pore proteins, water channels), ion and other pumps (e.g., calcium pumps, proton pumps), exchangers (e.g., sodium/potassium exchangers, sodium/hydrogen exchangers, potassium/hydrogen exchangers), electron transport proteins (e.g., cytochrome oxidase), enzymes and kinases (e.g., protein kinases, ATPases, GTPases, phosphatases, proteases.), structural/linker proteins (e.g., Caveolins, clathrin), adapter proteins (e.g., TRAD, TRAP, FAN), chemotactic/adhesion proteins (e.g., ICAM 1, selectins, CD34, VCAM-1, LFA-1, VLA-1), and chimeric/fusion proteins (e.g., proteins in which a normally soluble protein is attached to a transmembrane region of another protein).

[0674] Research products are designed for any specific type of application. These products may be packaged and distributed as, by way of non-limiting example, kits, chemicals, solutions, buffers, powders, solids, filters, columns, gels, matrixes, emulsions, pellets, capsules, and aerosols. Kits and reagents for certain research applications may be required by regulatory agency to be labeled "research use only" in order to indicate that the reagents are not intended for use in humans.

[0675] XII.B. Transfection

[0676] Transfection is the process of introducing genetic material into eukaryotic and archaeobacterial cells using biological, biochemical or physical methods. This process allows researchers to express and study target proteins in cultured cells (research use) as well as to deliver genetic material to cells in vivo or ex vivo systems (gene therapy). There are a variety of techniques which allow for the introduction and expression of proteins into target cells. These include mechanical transfection (Biolistic particles and Electroporation), calcium phosphate, DEAE-dextran/polybrene, viral based techniques and lipid based techniques.

[0677] The genetic material and/or nucleic acid to be delivered can be, by way of non-limiting example, nucleic acids that repair damaged or missing genes, nucleic acids for research applications, nucleic acids that kill a dysfunctional cell such as a cancer cell, antisense oligonucleotides to reduce or inhibit expression of a gene product, genetic material that increases expression of another gene, nucle-

otides and nucleotide analogs, peptide nucleic acids (PNAs), tRNAs, rRNAs, catalytic RNAs, RNA:DNA hybrid molecules, and combinations thereof.

[0678] The genetic material may comprise a gene expressing a protein. exemplary proteins include, but are not limited to, receptors (e.g., GPCRs, sphingolipid receptors, neurotransmitter receptors, sensory receptors, growth factor receptors, hormone receptors, chemokine receptors, cytokine receptors, immunological receptors, and compliment receptors, FC receptors), channels (e.g., potassium channels, sodium channels, calcium channels.), pores (e.g., nuclear pore proteins, water channels), ion and other pumps (e.g., calcium pumps, proton pumps), exchangers (e.g., sodium/potassium exchangers, sodium/hydrogen exchangers, potassium/hydrogen exchangers), electron transport proteins (e.g., cytochrome oxidase), enzymes and kinases (e.g., protein kinases, ATPases, GTPases, phosphatases, proteases), structural/linker proteins (e.g., Caveolins, clathrin), adapter proteins (e.g., TRAD, TRAP, FAN), chemotactic/adhesion proteins (e.g., ICAM 1, selectins, CD34, VCAM-1, LFA-1, VLA-1), and chimeric/fusion proteins (e.g., proteins in which a normally soluble protein is attached to a transmembrane region of another protein).

[0679] A minicell that is used to deliver therapeutic agents may comprise and display a binding moiety. By way of non-limiting example, binding moieties used for particular purposes may be a binding moiety directed to a compound or moiety displayed by a specific cell type or cells found predominantly in one type of tissue, which may be used, among other things, to target minicells and their contents to, specific cell types or tissues. A preferred binding moiety is an antibody or antibody derivative. Other binding moieties include, but are not limited to, receptors, enzymes, ligands, binding peptides, fusion proteins, small molecules conjugated to transmembrane proteins, ligands conjugated to transmembrane proteins, viral fusion proteins, and fusion/chimeric proteins.

[0680] A minicell containing genetic material may be to a target cell by methods including, but not limited to, receptor mediated endocytosis, cell fusion, or phagocytosis (Aderem et al., Mechanism of Phagocytosis in Macrophages, *Annu. Rev. Immunol.* 17:593-623, 1999). The minicell gene delivery system is used to deliver genetic material in culture for research applications as well as to cells in vivo as part of gene therapy or other therapeutic applications.

[0681] By way of non-limiting example, a minicell may express a protein such as invasins to induce receptor mediated endocytosis (Pepe et al., "*Yersinia enterocolitica* invasins: A primary role in the initiation of infection," *Proc. Natl. Acad. Sci. U.S.A.* 90:6473-6477, 1993; Alrutz et al., "Involvement of focal adhesion kinase in invasins-mediated uptake," *Proc. Natl. Acad. Sci. U.S.A.* 95:13658-13663, 1998). Invasins interact with the Beta2 Integrin protein and causes it to dimerize. Upon dimerization the Beta2 Integrin signals for an endocytotic event. Thus a minicell expressing the invasins protein will be taken up by cells expressing Beta2 Integrin via endocytosis.

[0682] Another non-limiting example of the minicell gene delivery and transfection system using invasins involves the expression of invasins following a targeting event. In this example, a minicell expresses a targeting protein that is capable of bringing the minicell in contact with a specific

target cell. Upon contact with the target cell, the minicell will be induced to transcribe and translate invasins. The induction is accomplished via signaling events or with a transcription factor dimerization event. The minicells can be engineered to contain targeting proteins that induce protein expression only upon contact with a specific target cell. By way of non-limiting example, the invasins is expressed only at the target cell where it induces endocytosis, thus preventing the minicell from entering any cell but the target cell.

[0683] Proteins can be induced and expressed post contact with target cells include but are not limited to antibodies and antibody derivatives, receptors, enzymes, ligands, binding peptides, fusion proteins, small molecules conjugated to transmembrane proteins, ligands conjugated to transmembrane proteins, viral fusion proteins, antibiotics, apoptotic proteins, hormones, toxins, poisons, and fusion/chimeric proteins.

[0684] Another non-limiting example of gene delivery or transfection using the minicell involves the use of the type III secretion apparatus of bacteria. The type III secretion apparatus is expressed in the minicell and used to transfer genetic material to a target cell.

[0685] Another non-limiting example of gene delivery and transfection using minicells involves minicells that have been engineered to contain anionic lipids or cationic lipids (Axel et al., "Toxicity, Uptake Kinetics and Efficacy of New Transfection Reagents: Increase of Oligonucleotide Uptake," *Jour. of Vasc. Res.* 040:1-14, 2000). Many types of lipids have been shown to induce or enhance transfection and gene delivery in a variety of cell types. Minicells containing such lipids could be used to transfer genetic material to specific cell types. Minicells can also be engineered to express targeting proteins that would allow the minicell to associate tightly with a target cell, which will facilitate the lipid interactions and gene transfer.

[0686] Another non-limiting example of gene delivery or transfection using minicells involves the use of ligands to induce receptor mediated endocytosis. By way of non-limiting example, the ligand is expressed on the surface of the minicell, or is attached to the surface of the minicell. A minicell containing genetic material is then able to associate with a target cell expressing the target receptor for the ligand. The receptor/ligand interaction will result in the endocytosis of the minicell into the target cell where the minicell would release and deliver the genetic material.

[0687] Another non-limiting example of gene delivery or transfection using minicells involves the use of fusion proteins, such as but not limited to viral capsid proteins. In this example the fusion protein would be expressed or attached to the outside of the minicell. The fusion protein would then induce fusion of a target cell with the minicell upon contact. The contact could be initiated via random non-targeting events or via the use of specific targeting proteins. In both cases the end result would be the fusion of the minicell with a target cell and the delivery of the genetic material.

[0688] XII.C. Non-Limiting Examples of Research Applications of Minicells

[0689] XII.C.1. Phage Interactions with Bacterial Membranes

[0690] One non-limiting example of a research application for minicells would be the study of phage interactions with a bacterial membrane. The minicells could be used to study

how phage associate and enter into a host bacterium. Another non-limiting example is the research application of minicells is to study isolated cell signaling pathways. The proteins of a signaling pathway could be expressed in the minicell and the signal cascade could be monitored. Another non-limiting example of research applications is the use of minicells to determine how recombination events occur. In this example the minicell is used to provide an environment to study the recombination event between two episomal plasmid DNA units.

**[0691]** XII.C.2. Matrices

**[0692]** Another non-limiting example of a research application of minicells is to form chromatography matrices for immunoprecipitation, isolation and separation techniques. The minicell can express and display target proteins with binding activity, including but not limited to antibodies and antibody derivatives. The minicell is then used to generate a matrix and loaded in a column or tube. The solution to be separated is mixed or passed through the column allowing the minicell to bind its target. The minicells are then separated away with the attached substance.

**[0693]** XII.C.3. Mutagenesis

**[0694]** Another non-limiting example of a research application for minicells involves site directed mutagenesis studies of target proteins. In this application minicells are generated to express target proteins with various mutations and deletions to study if function is compromised, enhanced or has an altered specificity for ligand binding.

**[0695]** XII.C.4. Metabolic Pathways

**[0696]** Another non-limiting example of research applications for minicells involves the study of metabolic rates of proteins and metabolites. The minicell can be generated to express metabolic pathways and the kinetics and function of that pathway can be studied.

**[0697]** XII.C.5. Cell Free Production of Proteins

**[0698]** Another non-limiting example of a research application for minicells involves uses in cell free production of functional proteins (Jermtus et al., Recent advances in producing and selecting functional proteins by using cell-free translation, Current Opinion in Biotechnology 9:534-548, 1999). Minicells can be prepared as a reagent used to prepare compositions for in vitro translation. As is described in detail elsewhere herein, the composition of minicells can be manipulated so as to be enriched for particular proteins or nucleic acids, including those involved in protein translation and folding and/or modification of the proteins so produced into functional forms, i.e., forms having the activity of the corresponding protein as it is isolated from natural sources. Non-limiting examples of such proteins and nucleic acids are ribosomal RNAs, ribosomal proteins, tRNAs, and the like.

**[0699]** XII.C.6. Assays

**[0700]** Minicells could also be used in manual, semi-automated, automated and/or robotic assays for the in vitro determinations of the compounds of interest including, by way of non-limiting example, ligands, proteins, small molecules, bioactive lipids, drugs, heavy metals, and the like in environmental samples (e.g., air, water, soil), blood, urine or tissue of humans or samples from non-human organisms

(e.g., plants, animals, protists) for the purpose of quantifying one or more compounds in a sample. A non-limiting example of this type of research applications is the expression on the surfaces of the minicells of a receptor such as the receptor that binds a toxin produced by *Bacillus anthracis*. The protein, protective antigen (PA), is a 82.7 kDa protein that binds one of the secreted anthrax toxins, lethal factor (LF) (see Price, B. et al., Infection and Immunity 69: 4509-4515. 2001). Minicells expressing the PA protein could be used to detect LF in an environmental sample or in human blood, urine or tissue for the purposes of determining the presence of anthrax. As a non-limiting example, a competitive binding assay or an antibody-based assay could be used to indicate binding of LF in the environmental or tissue sample. Another non-limiting example is the use of PA-expressing minicells in a lateral flow diagnostic where interaction between the minicells and the LF-containing sample is indicated by the presence of a colored reaction product on a test strip.

**[0701]** XIII. Minicell-Based Delivery of Biologically Active Agents

**[0702]** XIII.A. General Considerations

**[0703]** The minicells of the invention are capable of encapsulating and/or loading into a membrane a variety of substances, including but not limited to biologically active agents, including but not limited to diagnostic and therapeutic agents. Biologically active agents include, but are not limited to, nucleic acids, e.g., DNA, RNA, gene therapy constructs, ribozymes, antisense and other synthetic oligonucleotides including those with chemical modifications; peptide nucleic acids (PNAs); proteins; synthetic oligopeptides; peptomimetics; small molecules; radioisotopes; antibiotics; antibodies and antibody derivatives; and combinations and/or prodrugs of any of the preceding.

**[0704]** The surface of a minicell may be chemically altered in order to have certain properties that are desirable for their use as drug delivery agents. By way of non-limiting example, minicells may be chemically conjugated to polyethylene glycol (PEG), which provides for "stealth" minicells that are not taken as well and/or as quickly by the reticuloendothelial system (RES). Other compounds that may be attached to minicells include without limitation polysaccharides, polynucleotides, lipopolysaccharides, lipoproteins, glycosylated proteins, synthetic chemical compounds, and/or combinations of any of the preceding.

**[0705]** A minicell that is used to deliver therapeutic agents may comprise and display a binding moiety. By way of non-limiting example, binding moieties used for particular purposes may be a binding moiety directed to a compound or moiety displayed by a specific cell type or cells found predominantly in one type of tissue, which may be used, among other things, to target minicells and their contents to specific cell types or tissues. A preferred binding moiety is an antibody or antibody derivative, which are described in detail elsewhere herein. Other binding moieties include, but are not limited to, receptors, enzymes, ligands, binding peptides, fusion proteins, small molecules conjugated to transmembrane proteins, ligands conjugated to transmembrane proteins, viral fusion proteins, and fusion/chimeric proteins.

**[0706]** XIII.B. Cellular Uptake

**[0707]** In addition to binding moieties, proteins and other compounds that induce or enhance the uptake or fusion of the minicell with the target gene can be displayed on the surface of a minicell for applications involving the delivery of therapeutic agents, gene therapy, and/or transfection or other research applications. See, generally, *Adhesion Protein Protocols*, Vol. 96, Dejana, E. and Corada, M., eds., Humana Press, 1999.

**[0708]** XIII.B.1. Cellular Uptake Sequences from Eukaryotic Cells

**[0709]** Eukaryotic adhesion receptors, which mediate intercellular adhesion, can be used as agents or targets for cellular uptake: There are at least three distinct classes of adhesive molecules that leukocytes employ during their adhesive interactions (a) integrins, including but not limited to LEC-CAMS/Selectins (ELAM-1, LAM-1/Leu8/TQ1, and GMP140/PADGEM); (b) those belonging to the immunoglobulin superfamily including but not limited to CD2(LFA-2), CD3/TCR, CD4, CD8, CD28, CD44, CD54 (ICAM-1), ICAM-2, CD58 (LFA-3), VCAM-1, B7; and (c) Class I and II Major Histocompatibility Antigens (MHC).

**[0710]** The adhesion receptors that belong to the integrin family and control intercellular interactions are of particular interest. At least ten different structurally related cell surface heterodimeric (alpha and beta complexes) molecules have been defined as integrins and further classified into subfamilies (Springer T. A., 1990, *Nature* 346:425-434; Hynes, R. O., 1987, *Cell* 48:549-554; Moller, G. Editor, 1990, *Immunol. Rev.* 114:1-217). Each subfamily has a unique beta subunit, designated integrin beta1 (CD29), integrin beta2 (CD18), and integrin beta3 (CD61), each of which can associate with multiple alpha subunits, each with at least one di-valent cation binding site. The integrin family includes receptors for extracellular matrix components such as fibronectin, laminin, vitronectin, and collagen which recognize Arg-Gly-Asp in their ligands and utilize the beta1 or beta3 subunits (Springer T. A., 1990, *Nature* 346:425-434; Hynes, R. O., 1987, *Cell* 48:549-554; Hemler, M. E., 1988, *Immunol. Today* 9:109-113; Patarroyo, M., and Makgoba, M. W., 1989, *Scand. J. Immunol.* 30:129-164; Moller, G. Editor, 1990, *Immunol. Rev.* 114:1-217). There are at least six distinct alpha subunits alpha1 (CD49a), alpha2 (CD49b), alpha3 (CD49c), alpha4 (CD49d), alpha5 (CD49e), and alpha6 (CD49f) capable of associating with beta1 (CD29). The beta1 integrins are expressed on many nonhematopoietic and leukocyte cell types and are thought to play an active role in tissue organization by binding to extracellular matrix components found in many tissues and in the basement membranes underlying muscles, nervous system, epithelium and endothelium. While the expression of many beta1 integrins on leukocytes requires consistent activation, their expression on nonhematopoietic cells does not (Hemler, M. E., 1988, *Immunol. Today* 9:109-113; Patarroyo, M., and Makgoba, M. W., 1989, *Scand. J. Immunol.* 30:129-164). The complexity of the integrin family has been increased by the discovery of novel beta subunits beta3 (CD61), beta4 and beta5 that can associate with alpha 4, alpha 6, and alpha V subunits (Springer T. A., 1990, *Nature* 346:425-434; Hemler, M. E., 1988, *Immunol. Today* 9:109-113). This combinatorial use of alpha and beta subunits

confers considerable diversity in ligand recognition and also helps regulate communications between the inside and outside of the cell.

**[0711]** By way of non-limiting example, a minicell display an adhesion receptor, or a fusion protein that has a trans-membrane domain linked to a functional portion of an adhesion receptor. Such minicells will bind to cells displaying the ligand for the adhesion receptor.

**[0712]** XIII.B.2. Cellular Uptake Sequences from Prokaryotes

**[0713]** Bacterial adhesion proteins are another source of polypeptides that are used to stimulate uptake of minicells. See, generally, *Handbook of Bacterial Adhesion: Principles, Methods, and Applications*, Yuehwei H. An; Richard J. Friedman, eds., Humana Press, 2000; and Hultgren et al., "Bacterial Adhesions and Their Assembly," Chapter 150 in: *Escherichia coli and Salmonella typhimurium: Cellular and Molecular Biology*, 2<sup>nd</sup> Ed., Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1996, Volume 2, pages 1903-1999, and references cited therein.

**[0714]** By way of non-limiting example, a minicell may express a protein such as invasins to induce receptor mediated endocytosis (Pepe et al., *Yersinia enterocolitica* invasins: A primary role in the initiation of infection, *Proc. Natl. Acad. Sci. U.S.A.* 90:6473-6477, 1993; Alrutz et al., Involvement of focal adhesion kinase in invasins-mediated uptake, *Proc. Natl. Acad. Sci. U.S.A.* 95:13658-13663, 1998). Invasins interact with the Beta2 Integrin protein and causes it to dimerize. Upon dimerization the Beta2 Integrin signals for an endocytotic event. Thus a minicell expressing the invasins protein will be taken up by cells expressing Beta2 Integrin via endocytosis.

**[0715]** As another non-limiting example, the pneumococcal adhesin protein CpbA interacts with the human polyimmunoglobulin receptor (hpIgR) as either a part of the outer surface of a bacterial cell or as a free molecule Zhang et al. (*Cell* 102:827-837, 2000). The regions of CpbA:hpIgR interaction were mapped using a series of large peptide fragments derived from CpbA. CpbA (Swiss-Prot Accession No. 030874) contains a choline binding domain containing residues 454-663 and two N-terminal repetitive regions called R1 and R2 that are contained in residues 97-203 and 259-365, respectively. Polypeptides containing R1 and R2 interact with hpIgR, whereas polypeptides containing other sequences from CpbA do not bind to hpIgR. The R1 and/or R2 sequences of the CpbA polypeptide, and/or essentially identical, substantially identical, or homologous amino acid sequences, are used to facilitate the uptake of minicells by cells.

**[0716]** Another non-limiting example of gene delivery or transfection using the minicell involves the use of the type III secretion apparatus of bacteria. The type III secretion apparatus is expressed in the minicell and used to transfer genetic material to a target cell.

**[0717]** Other non-limiting examples of a minicell gene delivery and transfection targeting moiety are ETA (detoxified exotoxin a) protein delivery element described in U.S. Pat. No. 6,086,900 to Draper; Interalin and related proteins from *Listeria* species (Galan, *Alternative Strategies for Becoming an Insider: Lessons from the Bacterial World*,

Cell 103:363-366,2000); Intimin from pathogenic *E. coli* strains (Frankel et al., Intimin and the host cell—is it bound to end in Tir(s)? Trends in Microbiology 9:214-218); and SpeB, streptococcal pyrogenic exotoxin B (Stockbauer et al., A natural variant of the cysteine protease virulence factor of group A Streptococcus with an arginine-glycine-aspartic acid (RGD) motif preferentially binds human integrins  $\alpha_v\beta_3$  and  $\alpha_{IIb}\beta_3$  Proc. Natl. Acad. Sci. U.S.A. 96:242-247, 1999).

**[0718]** XIII.B.3. Cellular Uptake Sequences from Viruses

**[0719]** Cellular uptake sequences derived from viruses include, but are not limited to, the VP22 protein delivery element derived from herpes simplex virus-1 and vectors containing sequences encoding the VP22 protein delivery element are commercially available from Invitrogen (Carlsbad, Calif.; see also U.S. Pat. No. 6,017,735 to Ohare et al.); and the Tat protein delivery element derived from the amino acid sequence of the Tat protein of human immunodeficiency virus (HIV). See U.S. Pat. Nos. 5,804,604; 5,747,641; and 5,674,980.

**[0720]** XIII.B.4. Lipids

**[0721]** Another non-limiting example of gene delivery and transfection using minicells involves minicells that have been engineered to contain anionic lipids or cationic lipids (Axel et al., Toxicity, Uptake Kinetics and Efficacy of New Transfection Reagents: Increase of Oligonucleotide Uptake, Jour. of Vasc. Res. 040:1-14, 2000). Many types of lipids have been shown to induce or enhance transfection and gene delivery in a variety of cell types. Minicells containing such lipids could be used to transfer genetic material to specific cell types. Minicells can also be engineered to express targeting proteins that would allow the minicell to associate tightly with a target cell, which will facilitate the lipid interactions and gene transfer.

**[0722]** Another non-limiting example of gene delivery or transfection using minicells involves the use of ligands to induce receptor mediated endocytosis. By way of non-limiting example, the ligand is expressed on the surface of the minicell, or is attached to the surface of the minicell. A minicell containing genetic material is then able to associate with a target cell expressing the target receptor for the ligand. The receptor/ligand interaction will result in the endocytosis of the minicell into the target cell where the minicell would release and deliver the genetic material.

**[0723]** Another non-limiting example of gene delivery or transfection using minicells involves the use of fusion proteins, such as but not limited to viral capsid proteins. In this example the fusion protein would be expressed or attached to the outside of the minicell. The fusion protein would then induce fusion of a target cell with the minicell upon contact. The contact could be initiated via random non-targeting events or via the use of specific targeting proteins. In both cases the end result would be the fusion of the minicell with a target cell and the delivery of the genetic material.

**[0724]** XIII.C. Post-Targeting Expression of Cellular Uptake Sequences

**[0725]** Another non-limiting example of the minicell gene delivery and transfection system using invasin involves the expression of invasin following a targeting event. In this example, a minicell expresses a targeting protein that is

capable of bringing the minicell in contact with a specific target cell. Upon contact with the target cell, the minicell will be induced to transcribe and translate invasin. The induction is accomplished via signaling events or with a transcription factor dimerization event. The minicells can be engineered to contain targeting proteins that induce protein expression only upon contact with a specific target cell. By way of non-limiting example, the invasin is expressed only at the target cell where it induces endocytosis, thus preventing the minicell from entering any cell but the target cell.

**[0726]** Proteins can be induced and expressed post contact with target cells include but are not limited to antibodies and antibody derivatives, receptors, enzymes, ligands, binding peptides, fusion proteins, small molecules conjugated to transmembrane proteins, ligands conjugated to transmembrane proteins, viral fusion proteins, antibiotics, apoptotic proteins, hormones, toxins, poisons, and fusion/chimeric proteins.

**[0727]** XIII.D. Intracellular Targeting and Organellar Delivery

**[0728]** After delivery to and entry into a targeted cell, a minicell may be designed so as to be degraded, thereby releasing the therapeutic agent it encapsulates into the cytoplasm of the cell. The minicell and/or therapeutic agent may include one or more organellar delivery elements, which targets a protein into or out of a specific organelle or organelles. For example, the ricin A chain can be included in a fusion protein to mediate its delivery from the endosome into the cytosol. Additionally or alternatively, delivery elements for other organelles or subcellular spaces such as the nucleus, nucleolus, mitochondria, the Golgi apparatus, the endoplasmic reticulum (ER), the cytoplasm, etc. are included Mammalian expression constructs that incorporate organellar delivery elements are commercially available from Invitrogen (Carlsbad, Calif.: pShooter™ vectors). An H/KDEL (i.e., His/Lys-Asp-Glu-Leu sequence) may be incorporated into a fusion protein of the invention, preferably at the carboxy-terminus, in order to direct a fusion protein to the ER (see Andres et al., J. Biol. Chem. 266:14277-142782, 1991; and Pelham, Trends Bio. Sci. 15:483-486, 1990).

**[0729]** Another type of organellar delivery element is one which directs the fusion protein to the cell membrane and which may include a membrane-anchoring element. Depending on the nature of the anchoring element, it can be cleaved on the internal or external leaflet of the membrane, thereby delivering the fusion protein to the intracellular or extracellular compartment, respectively. For example, it has been demonstrated that mammalian proteins can be linked to i) myristic acid by an amide-linkage to an N-terminal glycine residue, to ii) a fatty acid or diacylglycerol through an amide- or thioether-linkage of an N-terminal cysteine, respectively, or covalently to iii) a phosphatidylinositol (PI) molecule through a C-terminal amino acid of a protein (for review, see Low, Biochem. J. 244: 1-13, 1987). In the latter case, the PI molecule is linked to the C-terminus of the protein through an intervening glycan structure, and the PI then embeds itself into the phospholipid bilayer; hence the term "GPI" anchor. Specific examples of proteins known to have GPI anchors and their C-terminal amino acid sequences have been reported (see Table 1 and FIG. 4 in Low, Biochemica et Biophysica Acta, 988:427-454, 1989;

and Table 3 in Ferguson, *Ann. Rev. Biochem.*, 57:285-320, 1988). Incorporation of GPI anchors and other membrane-targeting elements into the amino- or carboxy-terminus of a fusion protein can direct the chimeric molecule to the cell surface.

**[0730]** XIII.E. Minicell-Based Gene Therapy

**[0731]** The delivery of nucleic acids to treat diseases or disorders is known as gene therapy (Kay et al., *Gene Therapy*, *Proc. Natl. Acad. Sci. USA* 94:12744-12746, 1997). It has been proposed to use gene therapy to treat genetic disorders as well as pathogenic diseases. For reviews, see Desnick et al., *Gene Therapy for Genetic Diseases*, *Acta Paediatr. Jpn.* 40:191-203, 1998; and Bunnell et al., *Gene Therapy for Infectious Diseases*, *Clinical Microbiology Reviews* 11:42-56, 1998).

**[0732]** Gene delivery systems use vectors that contain or are attached to therapeutic nucleic acids. These vectors facilitate the uptake of the nucleic acid into the cell and may additionally help direct the nucleic acid to a preferred site of action, e.g., the nucleus or cytoplasm (Wu et al., "Delivery Systems for Gene Therapy," *Biotherapy* 3:87-95, 1991). Different gene delivery vectors vary with regards to various properties, and different properties are desirable depending on the intended use of such vectors. However, certain properties (for example, safety, ease of preparation, etc.) are generally desirable in most circumstances.

**[0733]** The minicells of the invention may be used as delivery agents for any therapeutic or diagnostic agent, including without limitation gene therapy constructs. Minicells that are used as delivery agents for gene therapy constructs may, but need not be, targeted to specific cells, tissues, organs or systems of an organism, of a pathogen thereof, using binding moieties as described in detail elsewhere herein.

**[0734]** In order to enhance the effectiveness of gene delivery vectors in, by way of non-limiting example, gene therapy and transfection, it is desirable in some applications of the invention to target specific cells or tissues of interest (targeted cells or tissues, respectively). This increases the effective dose (the amount of therapeutic nucleic acid present in the targeted cells or tissues) and minimizes side effects due to distribution of the therapeutic nucleic acid to other cells. For reviews, see Peng et al., "Viral Vector Targeting," *Curr. Opin. Biotechnol.* 10:454-457, 1999; Gunzburg et al., "Retroviral Vector Targeting for Gene Therapy," *Cytokines Mol. Ther.* 2:177-184, 1996.; Wickham, "Targeting Adenovirus," *Gene Ther.* 7:110-114, 2000; Dachs et al., "Targeting Gene Therapy to Cancer: A Review," *Oncol. Res.* 9:313-325, 1997; Curiel, "Strategies to Adapt Adenoviral Vectors for Targeted Delivery," *Ann NY Acad. Sci.* 886:158-171, 1999; Findeis et al., "Targeted Delivery of DNA for Gene Therapy via Receptors," *Trends Biotechnol.* 11:202-205, 1993.

**[0735]** Some targeting strategies make use of cellular receptors and their natural ligands in whole or in part. See, for example, Cristiano et al., "Strategies to Accomplish Gene Delivery Via the Receptor-Mediated Endocytosis Pathway," *Cancer Gene Ther.*, Vol. 3, No. 1, pp. 49-57, January-February 1996.; S. C. Phillips, "Receptor-Mediated DNA Delivery Approaches to Human Gene Therapy," *Biologicals*, Vol. 23, No. 1, pp. 13-6, March 1995; Michael et

al., "Strategies to Achieve Targeted Gene Delivery Via the Receptor-Mediated Endocytosis Pathway," *Gene Ther.*, Vol. 1, No. 4, pp. 223-32, July 1994; Lin et al., "Antiangiogenic Gene Therapy Targeting The Endothelium-Specific Receptor Tyrosine Kinase Tie2," *Proc. Natl. Acad. Sci., USA*, Vol. 95, pp. 8829-8834, 1998. Sudimack et al., "Targeted Drug Delivery Via the Folate Receptor," *Adv. Drug Deliv.*, pp. 147-62, March 2000; Fan et al., "Therapeutic Application of Anti-Growth Factor Receptor Antibodies," *Curr. Opin. Oncol.*, Vol. 10, No. 1, pp. 67-73, January 1998; Wadhwa et al., "Receptor Mediated Glycotargeting," *J. Drug Target*, Vol. 3, No. 2, pp. 111-27, 1995; Perales et al., "An Evaluation of Receptor-Mediated Gene Transfer Using Synthetic DNA-Ligand Complexes," *Eur. J. Biochem*, Vol. 1, No 2, pp. 226, 255-66, December 1994; Smith et al., "Hepatocyte-Directed Gene Delivery by Receptor-Mediated Endocytosis," *Semin Liver Dis.*, Vol. 19, No. 1, pp. 83-92, 1999.

**[0736]** Antibodies, particularly single-chain antibodies, to surface antigens specific for a particular cell type may also be used as targeting elements. See, for example, Kuroki et al., "Specific Targeting Strategies of Cancer Gene Therapy Using a Single Chain Variable Fragment (scFv) with a High Affinity for CEA," *Anticancer Res.*, pp. 4067-71, 2000; U.S. Pat. No. 6,146,885, to Dornburg, entitled "Cell-Type Specific Gene Transfer Using Retroviral Vectors Containing Antibody-Envelope Fusion Proteins"; Jiang et al., "In Vivo Cell Type-Specific Gene Delivery With Retroviral Vectors That Display Single Chain Antibodies," *Gene Ther.* 1999, 6:1982-7; Engelstadter et al., "Targeting Human T Cells By Retroviral Vectors Displaying Antibody Domains Selected From A Phage Display Library," *Hum. Gene Ther.* 2000, 11:293-303; Jiang et al., "Cell-Type-Specific Gene Transfer Into Human Cells With Retroviral Vectors That Display Single-Chain Antibodies," *J. Virol* 1998,72:10148-56; Chu et al., "Toward Highly Efficient Cell-Type-Specific Gene Transfer With Retroviral Vectors Displaying Single-Chain Antibodies," *J. Virol* 1997, 71:720-5; Chu et al., "Retroviral Vector Particles Displaying The Antigen-Binding Site Of An Antibody Enable Cell-Type-Specific Gene Transfer," *J. Virol* 1995, 69:2659-63; and Chu et al., "Cell Targeting With Retroviral Vector Particles Containing Antibody-Envelope Fusion Proteins," *Gene Ther.* 1994, 1:292-9.

**[0737]** Minicells are used to deliver DNA-based gene therapy to targeted cells and tissues. Double minicell transformants are used not only to target a particular cell/tissue type (e.g. HIV-infected T-cells) but are also engineered to fuse with and enter targeted cells and deliver a protein-based toxin (e.g., antibiotic, or pro-apoptotic gene like Bax), an antisense expression construct (e.g., antisense to a transcription factor), or antisense oligonucleotides (e.g., antisense to an anti-apoptotic gene such as Bcl-2. The doubly-transformed minicells express not only these cell death promoters, but also only target particular cells/tissues, thus minimizing toxicity and lack of specificity of gene therapy vectors. By "doubly-transformed" it is meant that the minicell comprises 2 expression elements, one eubacterial, the other eukaryotic. Alternatively, shuttle vectors, which comprise eubacterial and eukaryotic expression elements in one vector, may be used.

**[0738]** Minicell-based gene therapy is used to deliver expression plasmids that could correct protein expression deficiencies or abnormalities. As a non-limiting example, minicell inhalants are targeted to pulmonary alveolar cells



and are used to deliver chloride transporters that are deficient or otherwise material in cystic fibrosis. Protein hormone deficiencies (e.g., dwarfism) are corrected by minicell expression systems (e.g., growth hormone expression in pituitary cells). Duchene's muscular dystrophy is characterized by a mutation in the dystrophin gene; this condition is corrected by minicell-based gene therapy. Angiogenesis treatment for heart patients is made effective by FGF or VEGF-producing minicells targeted to the heart. In this case, plasmid-driven over-expression of these growth factors is preferred.

**[0739]** XIV. Therapeutic uses of Minicells

**[0740]** In addition to minicell-based gene therapy, minicells can be used in a variety of therapeutic modalities. Non-limiting examples of these modalities include the following applications.

**[0741]** XIV.A. Diseases and Disorders

**[0742]** Diseases and disorders to which the invention can be applied include, by way of non-limiting example, the following.

**[0743]** Diseases and disorders that involve the respiratory system, such as cystic fibrosis, lung cancer and tumors, asthma, pathogenic infections, allergy-related diseases and disorders, such as asthma; allergic bronchopulmonary aspergillosis; hypersensitivity pneumonia, eosinophilic pneumonia; emphysema; bronchitis; allergic bronchitis bronchiectasis; cystic fibrosis; hypersensitivity pneumonitis; occupational asthma; sarcoid, reactive airway disease syndrome, interstitial lung disease, hyper-eosinophilic syndrome, parasitic lung disease and lung cancer, asthma, adult respiratory distress syndrome, and the like;

**[0744]** Diseases and disorders of the digestive system, such as those of the gastrointestinal tract, including cancers, tumors, pathogenic infections, colitis; ulcerative colitis, diverticulitis, Crohn's disease, gastroenteritis, inflammatory bowel disease, bowel surgery ulceration of the duodenum, a mucosal villous disease including but not limited to coeliac disease, past infective villous atrophy and short gut syndromes, pancreatitis, disorders relating to gastrointestinal hormones, Crohn's disease, and the like;

**[0745]** Diseases and disorders of the skeletal system, such as spinal muscular atrophy, rheumatoid arthritis, osteoarthritis, osteoporosis, multiple myeloma-related bone disorder, cortical-striatal-spinal degeneration, and the like;

**[0746]** Autoimmune diseases, such as Rheumatoid arthritis (RA), multiple sclerosis (MS), Sjogren's syndrome, sarcoidosis, insulin dependent diabetes mellitus (IDDM), autoimmune thyroiditis, reactive arthritis, ankylosing spondylitis, scleroderma, polymyositis, dermatomyositis, psoriasis, vasculitis, Wegener's granulomatosis, Crohn's disease and ulcerative colitis amyotrophic lateral sclerosis, multiple sclerosis, autoimmune gastritis, systemic lupus erythematosus, autoimmune hemolytic anemia, autoimmune neutropenia, systemic lupus erythematosus, graft vs. host disease, bone marrow engraftment, some cases of Type I diabetes, and the like;

**[0747]** Neurological diseases and disorders, such as depression, bipolar disorder, schizophrenia, Alzheimer's disease, Parkinson's disease, familial tremors, Gilles de la Tourette syndrome, eating disorders, Lewy-body dementia, chronic pain and the like;

**[0748]** Pathological diseases and resultant disorders such as bacterial infections such as infection by *Escherichia*, *Shigella*, *Salmonella*; sepsis, septic shock, and bacteremia; infections by a virus such as HIV, adenovirus, smallpox virus, hepatovirus, and the like; and AIDS-related encephalitis, HIV-related encephalitis, chronic active hepatitis, and the like;

**[0749]** Proliferative disease and disorders, such as acute lymphoblastic leukemia, acute myelogenous leukemia, chronic myelogenous leukemia, metastatic melanoma, Kaposi's sarcoma, multiple myeloma, breast cancer, anal cancer, vulvar cancer, and the like; and

**[0750]** Various diseases, disorders and traumas including, but not limited to, apoptosis mediated diseases, inflammation, cerebral ischemia, myocardial ischemia, aging, sarcoidosis, granulomatous colitis, scleroderma, degenerative diseases, necrotic diseases, alopecia, neurological damage due to stroke, diffuse cerebral cortical atrophy, Pick disease, mesolimbocortical dementia, thalamic degeneration, Huntington chorea, cortical-basal ganglionic degeneration, cerebrotocerebellar degeneration, familial dementia with spastic paraparesis, polyglucosan body disease, Shy-Drager syndrome, olivopontocerebellar atrophy, progressive supranuclear palsy, dystonia musculorum deformans, Hallervorden-Spatz disease, Meige syndrome, acanthocytic chorea, Friedreich ataxia, Holmes familial cortical cerebellar atrophy, Gerstmann-Straussler-Scheinker disease, progressive spinal muscular atrophy, progressive bulbar palsy, primary lateral sclerosis, hereditary muscular atrophy, spastic paraplegia, glomerulonephritis, chronic thyroiditis, Grave's disease, thrombocytopenia, myasthenia gravis, psoriasis, peroneal muscular atrophy, hypertrophic interstitial polyneuropathy, hereditary ataxia polyneuritic form, optic neuropathy, and ophthalmoplegia.

**[0751]** A variety of diseases and disorders caused or exacerbated by pathogens may be treated using the invention. For a comprehensive description of pathogens and associated diseases and disorders, see Zinsser Microbiology, 20th Ed., Joklik, ed., Appelton-Century-Crofts, Norwalk, Conn., 1992, and references cited therein.

**[0752]** Minicells could also be used for replacement therapy (via gene therapy) in a variety of conditions known to be caused by protein or proteins that are either absent (e.g. Duchene's Muscular Dystrophy), reduced in level (Dwarfism) or aberrant (Sickle-cell anemia).

**[0753]** For a comprehensive description of diseases and disorders that may be treated using the invention, see *The Merck Manual of Diagnosis and Therapy*, 17th Ed., Beers et al., eds.; published edition, Merck and Co., Rahway, N.J., 1999; on-line edition, Medical Services, Usmedsa, USHH, <http://www.merck.com/pubs/mmanual/>, and references cited therein.

**[0754]** XIV.B. Removal of Toxins and Pathogens by Selective Absorption

**[0755]** When introduced into the bloodstream of an animal, receptor-displaying minicells bind and absorb toxic compounds, thereby removing such compounds from the general circulation. A therapeutic benefit ensues as the bound toxic compound cannot access the cells upon which it would otherwise exert its toxic effect.

[0756] Minicells expressing receptors for toxic substances are introduced IV in order to remove those toxins from the blood. One non-limiting example is in the treatment of sepsis. In one embodiment, a fusion protein is formed from the transmembrane domain of the EGF receptor or toxR and a known soluble receptor for LPS (lipopolysaccharide), such as the LBP (lipopolysaccharide binding protein) or the extracellular domain of CD 14 receptor protein, both of which bind the LPS bacterial endotoxin. These minicells inactivate LPS by initially binding to it and preventing LPS binding to naturally occurring CD14 receptors on heart cells and other cells involved in the endotoxic shock response. Eventually, the minicell-LPS complex is cleared from the blood by macrophages and other components of the immune system.

[0757] In another embodiment, minicells expressing receptors for toxic drugs (e.g., morphine) are used to treat drug overdoses. In other embodiments, minicells of the invention are used to express receptors to venoms (e.g., snake venom) or poisons (e.g., muscarinic receptor expression for the treatment of muscarine poisoning). In other embodiments, minicells of the invention expressing EDGRs are used to clear the blood of toxins and other undesirable compounds.

[0758] As another non-limiting example, minicells that bind pathogens are used to treat disease. Minicells, and pathogens bound thereto, may be ingested by human neutrophils and thus serve as an adjuvant to therapeutic processes mediated by neutrophils (Fox et al., Fate of the DNA in plasmid-containing *Escherichia coli* minicells ingested by human neutrophils, Blood 69:1394-400, 1987). In a related modality, minicells are used to bind compounds required for the growth of a pathogen.

#### [0759] XIV.C. Antiviral Therapy

[0760] In one modality, minicells of the invention are used as "sponges" for the selective absorption of any viral particle in the body. Without being limited to the following examples, minicells expression-receptors or antibodies selectively directed against viruses such as HIV, Hepatitis B and smallpox are used.

[0761] For the treatment of viremia, viruses are cleared from the blood by absorption during dialysis or by IV injection of minicells expressing targets for viral receptors. As the minicells interact with blood-borne virus particles, there is an initial reduction of host cell infection by virtue of the minicell-viral complexes that are formed. Since viral particles attach to and/or enter the minicell, they are not active because of the lack of machinery needed for viral replication in the minicells. The virus infected minicells are then cleared from the system by macrophages and processed by the immune system.

[0762] Certain retroviruses that infect particular host cells express viral proteins on the surfaces of the infected cells. HIV infection of T-cells is one non-limiting example of this. The viral protein, GP120, is expressed on the surfaces of infected T-cells (Turner et al., Structural Biology of HIV, J. Mol. Biol. 285:1-32, 1999). Minicells expressing CD4 act as anti-GP120 minicells not only to target virus particles in an infected patient, but also to identify infected T-cells. It may be desirable to also express co-receptors such as CCR5, CXCR4 or ARD (Dragic, An overview of the determinants of

CCR5 and CXCR4 co-receptor function, J. Gen. Virol. 82:1807-1814, 2001). The minicells are then used to kill the infected T-cells, or to inhibit viral replication and/or virion assembly.

[0763] In another non-limiting example of anti-pathogen therapy, minicells can be used to express bacterial surface antigens on their surfaces that facilitate cellular uptake of the minicell by intracellular pathogens such as *Mycobacterium tuberculosis* (the causative agent of tuberculosis), Rickettsiae, or viruses. In this "smart sponge" approach, selective absorption is accompanied by internalization of the pathogen by minicells. Destruction of the pathogen follows as a result of a combination of intraminicell digestion of pathogens and/or by the eventual processing of the virus-containing minicell by the cellular immune system of the patient.

#### [0764] XIV. D. Antibacterial and Antiparasitic Applications

[0765] Minicells may be used to kill pathogenic bacteria, protozoans, yeast and other fungi, parasitic worms, viruses and other pathogens by mechanisms that either do or do not rely on selective absorption. Antibiotics can be delivered to pathogenic organisms after first being targeted by the proteins or small molecules on the surfaces of the minicells that promote binding of the minicells to the surfaces of the pathogen. Fusion or injection of minicell contents into the pathogenic cell can result in the death or disablement of the pathogen and thus lower the effective dose of an antibiotic or gene therapeutic agent. Delivery of antibiotics tethered to or encapsulated by the minicells will reduce the effective dose of an antibiotic and will reduce its elimination by the renal system. In the case of delivering encapsulated molecules (e.g., antibiotics), purified/isolated minicells expressing membrane-bound proteins for targeting can be incubated with the molecules in vitro prior to administration. This would be particularly applicable to the use of protoplast minicells or poroplast minicells that have their outer membrane and cell wall or outer membrane only removed, respectively, thus facilitating the diffusion of the small molecule into the intact minicell.

[0766] Without being limited by the following example, minicells can be used as antibacterial agents by expressing on the surfaces of the minicells antigens, receptors, antibodies, or other targeting elements that will target the minicell to the pathogenic organism and facilitate the entry of plasmids, proteins, small molecules in order to gain access to or entry into the organism. Antibiotics may be encapsulated by minicells post isolation from the parent strain so that the antibiotic will not be effective against the minicell-producing bacteria itself. Since minicells are not able to reproduce, the antibiotic will not be lethal to the minicell delivery vehicle, but only to the targeted pathogen. In another non-limiting example, lysozymic factors e.g., complement may be expressed on the surfaces of the minicells or encapsulated by same as to promote lysis of the pathogen.

[0767] Minicells can also be engineered to express toxic proteins or other elements upon binding to the pathogen. Induction of minicell protein expression can be an event that is coincident with targeting or triggered by minicell binding to the target pathogen, thus making minicells toxic only when contact is made with the pathogenic organism. Minicells can be engineered to express fusion/chimeric proteins that are tethered to the membrane by transmembrane

domains that have signaling moieties on the cytoplasmic surfaces of the minicells, such as kinases or transcription factors. In one non-limiting example, a minicell fusion membrane-bound protein could be expressed containing an extracellular domain with a receptor, scFv, or other targeting protein that binds to the pathogen. The second segment of the chimera could be a transmembrane domain of a protein such as the EGF receptor or ToxR (that would tether the fusion protein to the membrane). Importantly, the cytoplasmic domain of the fusion protein could be a kinase that phosphorylates a bacterial transcription factor present in the minicell or could be fused to a transcription factor that would be expressed on the cytoplasmic surface of the minicell. The expression plasmid that was previously introduced into the minicells would then be activated by promoters utilizing the activated bacterial transcription factor pre-existing in the minicells or that which may be introduced by the minicell. Without being limited to the following example, the binding event could be signaled by a fusion protein containing elements of a receptor (e.g., EGF) or by an adhesion protein (e.g., an integrin) that require oligomerization. In the example of the use of integrins, bacterial or other transcription factors that also require dimerization could be cloned as fusion proteins such that the binding event would be signaled by a dimerization of two or more identical recombinant chimeric proteins that have association-dependent transcription factors tagged to the C-terminus of the fusion protein. The minicells would only be toxic when contact is made with the pathogen.

[0768] The proposed mechanism of induction coincident with targeting is not limited to the antiparasitic uses of minicells but can be used in other therapeutic situations where minicells are used to express proteins of therapeutic benefit when directed against eucaryotic cells of the organism (e.g., kill cancer cells with protein toxins expressed only after binding of the minicell to the cancer cell).

[0769] Transfer of DNA-containing plasmids or other expression element, antisense DNA, etc. may be used to express toxic proteins in the target cells or otherwise inhibit transcription and/or translation in the pathogenic organism or would otherwise be toxic to the cell. Without being limited by the following example, minicells can be used to transfer plasmids expressing growth repressors, DNAses, or other proteins or peptides (e.g., pro-apoptotic) that would be toxic to the pathogen.

#### [0770] XIV.E. Cancer Therapy

[0771] Fusion proteins expressed in minicells are used for cancer therapy. In a non-limiting example, phage display antibody libraries are used to clone single chain antibodies against tumor-associated (tumor-specific) antigens, such as MUC-1 or EGFvIII. Fusion proteins expressing these antibodies, and further comprising a single-pass transmembrane domain of an integral membrane protein, are used to "present" the antibody to the surface of the minicells. Injected minicells coated with anti-tumor antibodies target the tumor and deliver pro-apoptotic genes or other toxic substances to the tumor. The minicells are engulfed by the tumor cells by processes such receptor-mediated endocytosis (by, e.g., macrophages). By way of non-limiting example, toxR-invasin could be expressed on the surfaces of the minicells to promote endocytosis through the interaction between invasin and beta2-integrins on the surfaces of the target cells.

[0772] Fusion proteins possessing viral fusion-promoting proteins facilitate entry of the minicell to the tumor cell for gene therapy or for delivery of chemotherapy bioactive proteins and nucleic acids. In these and similar applications, the minicell may contain separate eukaryotic and eubacterial expression elements, or the expression elements may be combined into a single "shuttle vector."

#### [0773] XV. Diagnostic uses of Minicells

[0774] Minicells are transformed with plasmids expressing membrane-bound proteins, such as receptors, that bind to specific molecules in a particular biological sample such as blood, urine, feces, sweat, saliva or a tissue such as liver or heart. Minicells can also be used for delivery of therapeutic agents across the blood-brain barrier to the brain. This modality is used, by way of non-limiting example, for imaging purposes, and for the delivery of therapeutic agents, e.g., anti-depressants, and agents for the treatment of cancer, obesity, insomnia, schizophrenia, compulsive disorders and the like. Recombinant expression systems are incorporated into minicells where the plasmid-driven protein expression construct could be the produce a single gene product or a fusion protein, such as a soluble protein for the particular ligand fused with a transmembrane domain of a different gene. The fusion protein then acts as a membrane bound receptor for a particular ligand or molecule in the sample. Conventional cloning techniques (e.g., PCR) are used to identify genes for binding proteins, or phage display is used to identify a gene for a single-stranded variable antibody gene expressing binding protein for a particular ligand. The protein product is preferably a soluble protein. By constructing a plasmid containing this gene plus the transmembrane domain of a known single-pass membrane protein such as that of the EGF receptor, a fusion protein may be expressed on the surfaces of the minicells as an integral membrane protein with an extracellular domain that is preferably capable of binding ligand.

[0775] In another type of fusion protein, the transmembrane domain of the EGF receptor is fused to a known soluble receptor for a particular ligand, such as the LBP (lipopolysaccharide binding protein) or the extracellular domain of CD 14 receptor protein, both of which bind the bacterial endotoxin, LPS (lipopolysaccharide). The LBP/EGF or CD14/EGF fusion protein is used to measure LPS in the serum of patients suspected of sepsis.

[0776] The minicell system is used to express receptors such as those of the EDG (endothelial cell differentiation gene) family (e.g., EDG 1-9) that recognize sphingolipids such as sphingosine-1-phosphate (S1P), sphingosylphosphoryl choline (SPC) and the lysophospholipid, lysophosphatidic acid (LPA). Since these proteins are 7-pass integral membrane proteins, no additional transmembrane domains of another protein are needed, and the receptor protein is thus not a fusion protein.

[0777] Truncated or mutant forms of a protein of interest are useful in a diagnostic assay. For example, a protein that is an ligand-binding enzyme can be altered so as to bind its substrate of interest but can no longer convert substrate into product. One example of this application of minicell technology is the expression of a truncated or mutant lactic dehydrogenase which is able to bind lactic acid, but is not able to convert lactic acid to pyruvate. Similarly, hexokinase derivatives are used in minicells for glucose monitoring.

[0778] Minicells as diagnostic tools can be used either in vitro or in vivo. In the in vitro context, the minicells are used in an ELISA format or in a lateral flow diagnostic platform to detect the presence and level of a desired analyte. A sample (tissue, cell or body fluid sample) is taken and then tested in vitro. One advantage of the minicell system in detecting substances in tissue, cells or in body fluids is that the minicells can be used in vitro assays where the minicell is labeled with either a radioactive or fluorescent compound to aid in its detection in an ELISA format or lateral flow platform. Thus, the use of secondary antibody detection systems is obviated.

[0779] As an in vivo diagnostic, minicells can be radio-labeled. One method of labeling is to incubate minicells for a short time (about 8 hr) with a Ti/tracer (e.g., Tn99M) that is useful for detecting tumor metastases. The Tn99M accumulates in cells and loads into minicells after isolation or into the parent bacteria during growth phase. As Tn99M is oxidized by either the parent *E. coli* strain or by the minicells after isolation, the Tn99M is retained by the cell. Iodine-labeled proteins may also be used (Krown et al., TNF-alpha receptor expression in rat cardiac myocytes: TNF-alpha inhibition of L-type Ca<sup>2+</sup> transients, FEBS Letters 376:24-30, 1995).

[0780] One non-limiting example of in vivo detection of cancer making use of radiolabeled minicells is the use of the minicells to express chimeric membrane-bound single-chain antibodies against tumor-specific antigens (TSA) expressed on malignant melanoma or other transformed cells. Such TSAs include, but are not limited to, the breast cancer associated MUC1 antigen and variant forms of the EGFR (EGFvIII). By way of non-limiting example, minicells expressing antibodies to melanoma cells can be injected (IV) into a patient and then subjected to CAT scan of the lymphatic drainage in order to determine if a metastasis has occurred. This diagnostic technique obviates the need for lymph node dissection.

[0781] Another example of an in vivo diagnostic is to use the minicell system to express antibodies against oxidized low-density lipoproteins (LDL). Oxidized LDLs are associated with atherogenic plaques. Radiolabeled minicells (prepared as above) are injected IV into a person prior to nuclear imaging for image enhancement. MRI image contrast enhancement is performed by preparing minicells complexed (loaded) with contrast enhancers such as paramagnetic relaxivity agents and magnetic susceptibility agents.

[0782] In diagnostic as well as other applications, minicells preferentially detect a diagnostic marker, i.e., a marker associated with a disease or disorder. A diagnostic marker is statistically more likely to occur in individuals suffering from a disease than in those who are not diseased. Preferably, a diagnostic marker directly causes or is produced during a disease; however, the association may be no more than a correlation.

[0783] XVI. Drug Discovery (Screening) with Minicells

[0784] XVI.A. Assays

[0785] Minicells can be used in assays for screening pharmacological agents. By way of non-limiting example, the minicell system provides an environment for the expression of GPCRs and studies of their ligand binding kinetics. Such GPCR's include any member the Endothelial Differ-

entiation Gene (EDG) receptor family. GPCRs may participate in neoplastic cell proliferation, angiogenesis and cell death. Small molecules that either activate or inhibit the action of these GPCRs can be used in therapeutic interaction.

[0786] Assays are performed to determine protein expression and protein function. For example, the production of the protein can be followed using protein <sup>35</sup>S-Met labeling. This is performed by providing the cell only methionine that is labeled with <sup>35</sup>S. The cells are treated with IPTG to induce protein expression, and the <sup>35</sup>S-Met is incorporated into the protein. The cells are then lysed, and the resulting lysates were electrophoresed on an SDS gel and exposed to autoradiography film.

[0787] Another technique for assessing protein expression involves the use of western blots. Antibodies directed to various expressed proteins of interest have been generated and many are commercially available. Techniques for generating antibodies to proteins or polypeptides derived therefrom are known in the art (see, e.g., Cooper et al., Section III of Chapter 11 in: Short Protocols in Molecular Biology, 2nd Ed., Ausubel et al., eds., John Wiley and Sons, New York, 1992, pages 11-22 to 11-46). Standard western blot protocols, which may be used to show protein expression from the expression vectors in minicells and other expression systems, are known in the art. (see, e.g., Winston et al., Unit 10.7 of Chapter 10 in: Short Protocols in Molecular Biology, 2nd Ed., Ausubel et al., eds., John Wiley and Sons, New York, 1992, pages 10-32 to 10-35).

[0788] The amount of functional protein produced from a minicell expression system is determined through the use of binding studies. Ligands for the proteins of interest are used to show specific binding in the minicell system. Radiolabeled ligand is incubated with cells expressing the protein, in this case, a receptor for TNF-alpha. The cells are then centrifuged and the radioactivity counted in a scintillation counter. The amount of ligand that is bound reflects the amount of functional protein that is present in the sample.

[0789] By way of non-limiting example, the minicell system can be made to express EDGRs for the purpose of screening combinatorial chemistry libraries for molecules that enhance EDG activity. EDG activity is assayed in the minicell environment in several ways. One way is to crystallize minicells expressing an EDG protein (or any membrane-bound protein of choice) and then measure changes in the crystal structure to detect novel ligands. Circular dichroism (CD), x-ray diffraction, electron spin resonance (EPR) or other biophysical approaches are used to probe the structure of proteins in the minicell context. Additionally or alternately, minicells are produced that express not only the EDGR, but also express G-proteins (i.e., double transformants). An assay system involving GTP binding and hydrolysis is used to identify and assess which small molecules in the combinatorial chemistry library serve as activating ligands for EDG. The minicell expression system is used in in vitro binding assays and in high throughput drug screenings. The expression of mutant or truncated isoforms of proteins are used for functional analyses in order to discover inactive or overactive proteins for potential use in diagnostics or therapeutics.

[0790] XVI.B. High-Throughput Screening (HTS)

[0791] HTS typically uses automated assays to search through large numbers of compounds for a desired activity.

Typically HTS assays are used to find new drugs by screening for chemicals that act on a particular enzyme or molecule. For example, if a chemical inactivates an enzyme it might prove to be effective in preventing a process in a cell that causes a disease. High throughput methods enable researchers to try out thousands of different chemicals against each target very quickly using robotic handling systems and automated analysis of results.

[0792] As used herein, "high throughput screening" or "HTS" refers to the rapid *in vitro* screening of large numbers of compounds (libraries); generally tens to hundreds of thousands of compounds, using robotic screening assays. Ultra high-throughput Screening (uHTS) generally refers to the high-throughput screening accelerated to greater than 100,000 tests per day.

[0793] To achieve high-throughput screening, it is best to house samples on a multicontainer carrier or platform. A multicontainer carrier facilitates measuring reactions of a plurality of candidate compounds simultaneously. Multi-well microplates may be used as the carrier. Such multi-well microplates, and methods for their use in numerous assays, are both known in the art and commercially available.

[0794] Screening assays may include controls for purposes of calibration and confirmation of proper manipulation of the components of the assay. Blank wells that contain all of the reactants but no member of the chemical library are usually included. As another example, a known inhibitor (or activator) of an enzyme for which modulators are sought, can be incubated with one sample of the assay, and the resulting decrease (or increase) in the enzyme activity determined according to the methods herein. It will be appreciated that modulators can also be combined with the enzyme activators or inhibitors to find modulators which inhibit the enzyme activation or repression that is otherwise caused by the presence of the known enzyme modulator. Similarly, when ligands to a sphingolipid target are sought, known ligands of the target can be present in control/calibration assay wells.

[0795] The minicells of the invention are readily adaptable for use in high-throughput screening assays for screening candidate compounds to identify those which have a desired activity, e.g., inhibiting an enzyme that catalyzes a reaction that produces an undesirable compound, inhibiting function of a receptor independent of ligand interference, or blocking the binding of a ligand to a receptor therefor. The compounds thus identified can serve as conventional "lead compounds" or can themselves be used as therapeutic agents.

[0796] The methods of screening of the invention comprise using screening assays to identify, from a library of diverse molecules, one or more compounds' having a desired activity. A "screening assay" is a selective assay designed to identify, isolate, and/or determine the structure of, compounds within a collection that have a preselected activity. By "identifying" it is meant that a compound having a desirable activity is isolated, its chemical structure is determined (including without limitation determining the nucleotide and amino acid sequences of nucleic acids and polypeptides, respectively) the structure of and, additionally or alternatively, purifying compounds having the screened activity). Biochemical and biological assays are designed to test for activity in a broad range of systems ranging from

protein-protein interactions, enzyme catalysis, small molecule-protein binding, agonists and antagonists, to cellular functions. Such assays include automated, semi-automated assays and HTS (high throughput screening) assays.

[0797] In HTS methods, many discrete compounds are preferably tested in parallel by robotic, automatic or semi-automatic methods so that large numbers of test compounds are screened for a desired activity simultaneously or nearly simultaneously. It is possible to assay and screen up to about 6,000 to 20,000, and even up to about 100,000 to 1,000,000 different compounds a day using the integrated systems of the invention.

[0798] Typically in HTS, target molecules are contained in each well of a multi-well microplate; in the case of enzymes, reactants are also present in the wells. Currently, the most widely established techniques utilize 96-well microtiter plates. In this format, 96 independent tests are performed simultaneously on a single 8 cm x 12 cm plastic plate that contains 96 reaction wells. One or more blank wells contains all of the reactants except the candidate compound. Each of the non-standard wells contain at least one candidate compound.

[0799] These wells typically require assay volumes that range from 50 to 500  $\mu$ l. In addition to the plates, many instruments, materials, pipettors, robotics, plate washers and plate readers are commercially available to fit the 96-well format to a wide range of homogeneous and heterogeneous assays. Microtiter plates with more wells, such as 384-well microtiter plates, are also used, as are emerging methods such as the nanowell method described by Schullek et al. (*Anal Biochem.*, 30 246, 20-29, 1997).

[0800] In one modality, screening comprises contacting a sphingolipid target with a diverse library of member compounds, some of which are ligands of the target, under conditions where complexes between the target and ligands can form, and identifying which members of the libraries are present in such complexes. In another non limiting modality, screening comprises contacting a target enzyme with a diverse library of member compounds, some of which are inhibitors (or activators) of the target, under conditions where a product or a reactant of the reaction catalyzed by the enzyme produce a detectable signal. In the latter modality, inhibitors of target enzyme decrease the signal from a detectable product or increase a signal from a detectable reactant (or vice-versa for activators).

[0801] Minicells of the invention expressing and/or displaying a protein are used for screening assays designed to identify agents that modulate the activity of the target protein. Such assays include competitive inhibition binding assays for high throughput assays. Competitive inhibition assays include but are not limited to assays that screen agents against a specific target protein to identify agents that inhibit, interfere, block, or compete with protein-ligand interactions, protein-protein interactions, enzymatic activity, or function of a specific protein. Examples of competitive inhibition include but are not limited to the development of neutral inhibitors of the serine protease factor Xa that were discovered using a high throughput screening assay using a compound library (Carr et al, *Neutral inhibitors of the serine protease factor Xa*, *Bioorg Med Chem Lett* 11, 2001), the design and characterization of potent inhibitors for the human oxytocin receptor (Seyer et al, *Design, synthesis and*

pharmacological characterization of a potent radio iodinated and photoactivatable peptidic oxytocin antagonist, *J Med Chem.* 44:3022-30, 2001), and the identification of non-peptide somatostatin antagonists of the sst(3) protein (Thuriereau et al, Identification of potent non-peptide somatostatin antagonists with sst(3) selectivity, *J Med Chem.* 44:2990-3000, 2001).

**[0802]** High throughput competitive inhibition assays are designed to identify agents that inhibit a specific target protein. Such assays include but are not limited to ones that measure enzymatic activity, protein-ligand interactions, protein-protein interactions and other functions of proteins. Minicells that express and/or display a specific protein could be used in all types of competitive inhibition assays.

**[0803]** One non-limiting example of high throughput competitive inhibition screening using minicells for the purpose of this patent involves the competitive inhibition of known ligands. The ligand is attached to but not limited to a fluorophore, fluorescent protein, tags such as 6x His tag or FLAG tag, chromophores, radiolabeled proteins and molecules, binding moieties such as avidin and streptavidin, voltage sensitive dyes and proteins, bioluminescent proteins and molecules, or fluorescent peptides. The target protein, which binds the tagged ligand, is expressed and stably displayed by the minicell. When the ligand is added to the minicell solution the ligand binds to the target protein. Following a wash the interaction is detected via the fluorophore, fluorescent protein, tag, or fluorescent peptide. The ligand-bound minicells could either be centrifuged (taking advantage of the sedimentation properties of the minicell particle) or immunoprecipitated with an antibody against an antigen expressed on the minicell membrane or the minicells can be adsorbed/fixated to a substrate such as a standard 96 well plate. The competitive inhibition assay is carried out by adding agents to the minicell mix either before, together or after the ligand is added. Thus if the agent is a competitive inhibitor of the ligand to the target protein the ligand will be washed away from the minicell because it is not associated with the target protein. The agent prevents binding and thus eliminated the detection signal from the minicell.

**[0804]** Minicells of this invention are used in "functional screening HTS assays". Functional screening assays are defined as assays that provide information about the function of a specific target protein. Functional assays screen agents against specific target proteins to identify agents that either act as antagonist or as an agonist against the protein. Functional assays require that the target protein be in an environment that allows it to carry out its natural function. Such functions include but are not limited to G-proteins coupling with a GPCR, enzymatic activity such as phosphorylation or proteolysis, protein-protein interaction, and transport of molecules and ions.

**[0805]** Functional assays screen agents against proteins which are capable of natural function. Target proteins used in functional studies must carry out a function that is measurable. Examples of protein functions that are measurable include but are not limited to the use of Fluorescent Resonance Energy Transfer (FRET) to measure the G-protein coupling to a GPCR (Ruiz-Velasco et al., Functional expression and FRET analysis of green fluorescent proteins fused to G-protein subunits in rat sympathetic neurons, *J. Physiol.* 537:679-692, 2001; Janetopoulos et al., Receptor-

mediated activation of heterotrimeric G-proteins in living cells, *Science* 291:2408-2411, 2001); Bioluminescence Resonance Energy Transfer (BRET) to assay for functional ligand induced G-protein coupling to a target GPCR (Menard, L. Bioluminescence Resonance Energy Transfer (BRET): A powerful platform to study G-protein coupled receptors (GPCR) activity in intact cells, *Assay Development*, Nov. 28-30, 2001), the use of fluorescent substrates to measure the enzymatic activity of proteases (Grant, Designing biochemical assays for proteases using fluorogenic substrates, *Assay Development*, Nov. 28-30, 2001); and the determination of ion channel function via the use of voltage sensitive dyes (Andrews et al, Correlated measurements of free and total intracellular calcium concentration in central nervous system neurons, *Microsc Res Tech.* 46:370-379, 1999).

**[0806]** One non-limiting example of high throughput functional screening assay using minicells for the purpose of this patent involves the functional coupling of GPCRs to their respective G-protein. Upon ligand binding, voltage polarization, ion binding, light interaction and other stimulatory events activate GPCRs and cause them to couple to their respective G-protein. In a minicell, both the GPCR and its respective G-proteins can be simultaneously expressed. Upon activation of the GPCR the coupling event will occur in the minicell. Thus by detecting this coupling in the minicell, one could screen for agents that bind GPCRs to identify antagonists and agonists. The antagonists are identified using inhibition assays that detect the inhibition of function of the GPCR. Thus the agent interacts with the GPCR in a way that it inhibits the GPCR from being activated. The agonists are identified by screening for agents that activate the GPCR in the absence of the natural activator.

**[0807]** The detection of GPCR activation and coupling in a minicell is accomplished by using systems that generate a signal upon coupling. One non-limiting example involves the use of BRET or FRET. These systems require that two fluorescent or bioluminescent molecules or proteins be brought into close contact. Thus by attaching one of these molecules or proteins to the GPCR and one to the G-protein, they will be brought together upon coupling and a signal will be generated. This signal can be detected using specific detection equipment and the coupling event can be monitored. Thus the function of the GPCR can be assayed and used in functional assays in minicells.

**[0808]** Another non-limiting functional assay for GPCRs and other proteins in minicells involves the use of transcription factors. Many bacterial transcription factors and eukaryotic transcription factors require dimerization for activation. By attaching one subunit of a transcription factor to a GPCR and the other subunit to a G-protein, the subunits will dimerize upon coupling of the GPCR to the G-protein because they will be brought into close contact. The dimerized transcription factor will then be activated and will act on its target episomal DNA. In the minicell system the episomal DNA target will be a plasmid that encodes for proteins that provide a signal for detection. Such proteins include but are not limited to luciferase; green fluorescent protein (GFP), and derivatives thereof such as YFP, BFP, etc.; alcohol dehydrogenase, and other proteins that can be assayed for expression. The activation of the GPCR will result in coupling and activation of the transcription factor.

The transcription factor will then induce transcription and translation of specific detector proteins. Thus the activation of the GPCR will be monitored via the expression of the detector protein.

[0809] In another modality, the transcription factor can inhibit expression in the minicell system and thus allowing for the screening of constitutively active GPCRs and proteins. For example if the GPCR were constitutively active then the transcription factor to use would be one that inhibits transcription and translation. Thus agents could be screened against the constitutively active GPCR to identify agents that caused the constitutively active GPCR to uncouple. The uncoupling will result in the inactivation of the transcription factor. The inhibition caused by the transcription factor will be removed and transcription and translation will occur. Thus a detectable protein will be made and a signal will be received.

[0810] The transcription dimerization assay can be used for any protein function that involves a protein-protein interaction, protein-ligand interaction and protein-drug interaction. Thus any assay involving such interactions can be carried out in the minicell.

[0811] Another non-limiting functional screening assay involves the use of enzymatic function to screen for functionality. In this modality the receptor or other protein performs a specific enzymatic function. This function is then carried out in the minicell and monitored using biochemical and other techniques. For example if the target protein was a protease then fluorescent peptides with the cleavage site of the protease could be used to monitor the activity of the protease. If the protease was functioning then the peptide would be cleaved and the fluorescents would change. Thus agents can be screened against the protease in the minicell system and the fluorescents can be monitored using specific detection systems. In another non-limiting example, a membrane-bound enzyme such as sphingomyelinase could be expressed in minicells and the minicell particles adsorbed to a standard substrate such as a 96 well plate. The enzymatic activity could be assessed by a standard in vitro assay involving the release of product (phosphocholine) (e.g., Amplex™ kit A-12220 sold by Molecular Probes). Sphingomyelinase inhibitors could be screened by measuring the reduction of phosphocholine production in the well when presented with substrate (sphingomyelin) in a coupled fluorescence assay.

[0812] Another non-limiting example of minicells used for functional assays involves the screening of agonists/antagonists for ion channels. In this example the calcium channel, SCaMPER, is encoded on a polycistronic episomal plasmid, which also encodes for a luminescent soluble protein, aequorin. In this assay, the minicell will contain aequorin proteins in its cytoplasm and SCaMPER proteins expressed on the minicell membrane. Thus upon activation of SCaMPER by its ligand, SPC, or by an analog thereof, calcium will flow into the minicell and will be bound by the aequorin which will luminescence. Thus a detection signal for the functional activation of the calcium channel is obtained.

[0813] Minicell can also be employed for expression of target proteins and the preparation of membrane preparations for use in screening assays. Such proteins include but are not limited to receptors (e.g., GPCRs, sphingolipid

receptors, neurotransmitter receptors, sensory receptors, growth factor receptors, hormone receptors, chemokine receptors, cytokine receptors, immunological receptors, and compliment receptors, FC receptors), channels (e.g., potassium channels, sodium channels, calcium channels.), pores (e.g., nuclear pore proteins, water channels), ion and other pumps (e.g., calcium pumps, proton pumps), exchangers (e.g., sodium/potassium exchangers, sodium/hydrogen exchangers, potassium/hydrogen exchangers), electron transport proteins (e.g., cytochrome oxidase), enzymes and kinases (e.g., protein kinases, ATPases, GTPases, phosphatases, proteases.), structural/linker proteins (e.g., Caveolins, clathrin), adapter proteins (e.g., TRAD, TRAP, FAN), chemotactic/adhesion proteins (e.g., ICAM11, selectins, CD34, VCAM-1, LFA-1, VLA-1), and chimeric/fusion proteins (e.g., proteins in which a normally soluble protein is attached to a transmembrane region of another protein). In such assays the membrane preparations are used to screen for agents that are either antagonists or agonists. These assays use various formats including but not limited to competitive inhibition.

[0814] The format for the screening of minicells includes but is not limited to the use of test tubes, 6 well plates, 12 well plates, 24 well plates, 96 well plates, 384 well plates, 1536 well plates, and other microtiter well plates. In these systems the minicells can be immobilized, attached, bound, or fused with the above test tubes or plates. The minicells can also be free in solution for use in tubes and plates. The detection systems for the minicell assay include but are not limited to fluorescent plate readers, scintillation counters, spectrophotometers, Viewlux CCD Imager, Luminex, ALPHAQuest, BIAcore, FLIPR and F-MAT. Minicell assays can be carried out with but not limited to techniques such as manual handling, liquid handlers, robotic automated systems and other formats.

[0815] XVI.C. Chemical Libraries

[0816] Developments in combinatorial chemistry allow the rapid and economical synthesis of hundreds to thousands of discrete compounds. These compounds are typically arrayed in moderate-sized libraries of small organic molecules designed for efficient screening. Combinatorial methods, can be used to generate unbiased libraries suitable for the identification of novel inhibitors. In addition, smaller, less diverse libraries can be generated that are descended from a single parent compound with a previously determined biological activity. In either case, the lack of efficient screening systems to specifically target therapeutically relevant biological molecules produced by combinatorial chemistry such as inhibitors of important enzymes hampers the optimal use of these resources.

[0817] A combinatorial chemical library is a collection of diverse chemical compounds generated by either chemical synthesis or biological synthesis, by combining a number of chemical "building blocks," such as reagents. For example, a linear combinatorial chemical library, such as a polypeptide library, is formed by combining a set of chemical building blocks (amino acids) in a large number of combinations, and potentially in every possible way, for a given compound length (i.e., the number of amino acids in a polypeptide compound). Millions of chemical compounds can be synthesized through such combinatorial mixing of chemical building blocks.

[0818] A "library" may comprise from 2 to 50,000,000 diverse member compounds. Preferably, a library comprises at least 48 diverse compounds, preferably 96 or more diverse compounds, more preferably 384 or more diverse compounds, more preferably, 10,000 or more diverse compounds, preferably more than 100,000 diverse members and most preferably more than 1,000,000 diverse member compounds. By "diverse" it is meant that greater than 50% of the compounds in a library have chemical structures that are not identical to any other member of the library. Preferably, greater than 75% of the compounds in a library have chemical structures that are not identical to any other member of the collection, more preferably greater than 90% and most preferably greater than about 99%.

[0819] The preparation of combinatorial chemical libraries is well known to those of skill in the art. For reviews, see Thompson et al., Synthesis and application of small molecule libraries, *Chem Rev* 96:555-600, 1996; Kenan et al., Exploring molecular diversity with combinatorial shape libraries, *Trends Biochem Sci* 19:57-64, 1994; Janda, Tagged versus untagged libraries: methods for the generation and screening of combinatorial chemical libraries, *Proc Natl Acad Sci USA*. 91:10779-85, 1994; Lebl et al., One-bead-one-structure combinatorial libraries, *Biopolymers* 37:177-98, 1995; Eichler et al., Peptide, peptidomimetic, and organic synthetic combinatorial libraries, *Med Res Rev*. 15:481-96, 1995; Chabala, Solid-phase combinatorial chemistry and novel tagging methods for identifying leads, *Curr Opin Biotechnol*. 6:632-9, 1995; Dolle, Discovery of enzyme inhibitors through combinatorial chemistry, *Mol Divers*. 2:223-36, 1997; Fauchere et al., Peptide and non-peptide lead discovery using robotically synthesized soluble libraries, *Can J Physiol Pharmacol*. 75:683-9, 1997; Eichler et al., Generation and utilization of synthetic combinatorial libraries, *Mol Med Today* 1: 174-80, 1995; and Kay et al., Identification of enzyme inhibitors from phage-displayed combinatorial peptide libraries, *Comb Chem High Throughput Screen* 4:535-43, 2001.

[0820] Such combinatorial chemical libraries include, but are not limited to, peptide libraries (see, e.g., U.S. Pat. No. 5,010,175, Furka, *Int. J. Pept. Prot. Res.*, 37:487-493 (1991) and Houghton, et al., *Nature*, 354:84-88 (1991)). Other chemistries for generating chemical diversity libraries can also be used. Such chemistries include, but are not limited to, peptoids (PCT Publication No. WO 91/19735); encoded peptides (PCT Publication WO 93/20242); random bioligomers (PCT Publication No. WO 92/00091); benzodiazepines (U.S. Pat. No. 5,288,514); diversomers, such as hydantoins, benzodiazepines and dipeptides (Hobbs, et al., *Proc. Nat. Acad. Sci. USA*, 90:6909-6913 (1993)); vinylogous polypeptides (Hagihara, et al., *J. Amer. Chem. Soc.* 114:6568 (1992)); nonpeptidic peptidomimetics with  $\beta$ -D-glucose scaffolding (Hirschmann, et al., *J. Amer. Chem. Soc.*, 114:9217-9218 (1992)); analogous organic syntheses of small compound libraries (Chen, et al., *J. Amer. Chem. Soc.*, 116:2661 (1994)); oligocarbamates (Cho, et al., *Science*, 261:1303 (1993)); and/or peptidyl phosphonates (Campbell, et al., *J. Org. Chem.* 59:658 (1994)); nucleic acid libraries (see, Ausubel, Berger and Sambrook, all supra); peptide nucleic acid libraries (see, e.g., U.S. Pat. No. 5,539,083); antibody libraries (see, e.g., Vaughn, et al., *Nature Biotechnology*, 14(3):309-314 (1996) and PCT/US96/10287); carbohydrate libraries (see, e.g., Liang, et al., *Science*, 274:1520-1522 (1996) and U.S. Pat. No. 5,593,853); small

organic molecule libraries (see, e.g., benzodiazepines, *Baum C&E News*, January 18, page 33 (1993); isoprenoids (U.S. Pat. No. 5,569,588); thiazolidinones and metathiazanones (U.S. Pat. No. 5,549,974); pyrrolidines (U.S. Pat. Nos. 5,525,735 and 5,519,134); morpholino compounds (U.S. Pat. No. 5,506,337); benzodiazepines (U.S. Pat. No. 5,288,514); and the like.

[0821] Devices for the preparation of combinatorial libraries are commercially available (see, e.g., 357 MPS, 390 MPS, Advanced Chem. Tech, Louisville Ky., Symphony, Rainin, Woburn, Mass., 433A Applied Biosystems, Foster City, Calif., 9050 Plus, Millipore, Bedford, Mass.). In addition, numerous combinatorial libraries are themselves commercially available (see, e.g., ComGenex, Princeton, N.J., Asinex, Moscow, Ru, Tripos, Inc., St. Louis, Mo., Chem-Star, Ltd., Moscow, RU, 3D Pharmaceuticals, Exton, Pa., Martek Bio sciences, Columbia, Md., etc.).

[0822] XVI.D. Measuring Enzymatic and Binding Reactions During Screening Assays

[0823] Techniques for measuring the progression of enzymatic and binding reactions in multicontainer carriers are known in the art and include, but are not limited to, the following.

[0824] Spectrophotometric and spectrofluorometric assays are well known in the art. Examples of such assays include the use of colorimetric assays for the detection of peroxides, as disclosed in Example 1(b) and Gordon, A. J. and Ford, R. A., *The Chemist's Companion: A Handbook Of Practical Data, Techniques, And References*, John Wiley and Sons, N.Y., 1972, Page 437.

[0825] Fluorescence spectrometry may be used to monitor the generation of reaction products. Fluorescence methodology is generally more sensitive than the absorption methodology. The use of fluorescent probes is well known to those skilled in the art. For reviews, see Bashford et al., *Spectrophotometry and Spectrofluorometry: A Practical Approach*, pp. 91-114, IRL Press Ltd. (1987); and Bell, *Spectroscopy In Biochemistry*, Vol. I, pp. 155-194, CRC Press (1981).

[0826] In spectrofluorometric methods, enzymes are exposed to substrates that change their intrinsic fluorescence when processed by the target enzyme. Typically, the substrate is nonfluorescent and converted to a fluorophore through one or more reactions. As a non-limiting example, SMase activity can be detected using the Amplex® Red reagent (Molecular Probes, Eugene, Oreg.). In order to measure sphingomyelinase activity using Amplex Red, the following reactions occur. First, SMase hydrolyzes sphingomyelin to yield ceramide and phosphorylcholine. Second, alkaline phosphatase hydrolyzes phosphorylcholine to yield choline. Third, choline is oxidized by choline oxidase to betaine. Finally,  $H_2O_2$ , in the presence of horseradish peroxidase, reacts with Amplex Red to produce the fluorescent product, Resorufin, and the signal therefrom is detected using spectrofluorometry.

[0827] Fluorescence polarization (FP) is based on a decrease in the speed of molecular rotation of a fluorophore that occurs upon binding to a larger molecule, such as a receptor protein, allowing for polarized fluorescent emission by the bound ligand. FP is empirically determined by measuring the vertical and horizontal components of fluo-



rophore emission following excitation with plane polarized light. Polarized emission is increased when the molecular rotation of a fluorophore is reduced. A fluorophore produces a larger polarized signal when it is bound to a larger molecule (i.e. a receptor), slowing molecular rotation of the fluorophore. The magnitude of the polarized signal relates quantitatively to the extent of fluorescent ligand binding. Accordingly, polarization of the "bound" signal depends on maintenance of high affinity binding.

[0828] FP is a homogeneous technology and reactions are very rapid, taking seconds to minutes to reach equilibrium. The reagents are stable, and large batches may be prepared, resulting in high reproducibility. Because of these properties, FP has proven to be highly automatable, often performed with a single incubation with a single, premixed, tracer-receptor reagent. For a review, see Owickiet al., *Application of Fluorescence Polarization Assays in High-Throughput Screening*, Genetic Engineering News, 17:27, 1997.

[0829] FP is particularly desirable since its readout is independent of the emission intensity (Checovich, W. J., et al., *Nature* 375:254-256, 1995; Dandliker, W. B., et al., *Methods in Enzymology* 74:3-28, 1981) and is thus insensitive to the presence of colored compounds that quench fluorescence emission. FP and FRET (see below) are well-suited for identifying compounds that block interactions between receptors and their ligands. See, for example, Parker et al., *Development of high throughput screening assays using fluorescence polarization: nuclear receptor-ligand-binding and kinase/phosphatase assays*, *J Biomol Screen* 5:77-88, 2000.

[0830] Exemplary normal-and-polarized fluorescence readers include the POLARION fluorescence polarization system (Tecan A G, Hombrechtikon, Switzerland). General multiwell plate readers for other assays are available, such as the VERSAMAX reader and the SPECTRAMAX multiwell plate spectrophotometer (both from Molecular Devices).

[0831] Fluorescence resonance energy transfer (FRET) is another useful assay for detecting interaction and has been described previously. See, e.g., Heim et al., *Curr. Biol.* 6:178-182, 1996; Mitra et al., *Gene* 173:13-17 1996; and Selvin et al., *Meth. Enzymol.* 246:300-345, 1995. FRET detects the transfer of energy between two fluorescent substances in close proximity, having known excitation and emission wavelengths. As an example, a protein can be expressed as a fusion protein with green fluorescent protein (GFP). When two fluorescent proteins are in proximity, such as when a protein specifically interacts with a target molecule, the resonance energy can be transferred from one excited molecule to the other. As a result, the emission spectrum of the sample shifts, which can be measured by a fluorometer, such as a FMAX multiwell fluorometer (Molecular Devices, Sunnyvale Calif.).

[0832] Scintillation proximity assay (SPA) is a particularly useful assay for detecting an interaction with the target molecule. SPA is widely used in the pharmaceutical industry and has been described (Hanselman et al., *J. Lipid Res.* 38:2365-2373 (1997); Kahl et al., *Anal. Biochem.* 243:282-283 (1996); Udenfried et al., *Anal. Biochem.* 161:494-500 (1987)). See also U.S. Pat. Nos. 4,626,513 and 4,568,649, and European Patent No. 0,154,734. An exemplary com-

mercially available system uses FLASHPLATE scintillant-coated plates (NEN Life Science Products, Boston, Mass.).

[0833] The target molecule can be bound to the scintillator plates by a variety of well known means. Scintillant plates are available that are derivatized to bind to fusion proteins such as GST, His6 or Flag fusion proteins. Where the target molecule is a protein complex or a multimer, one protein or subunit can be attached to the plate first, then the other components of the complex added later under binding conditions, resulting in a bound complex.

[0834] In a typical SPA assay, the gene products in the expression pool will have been radiolabeled and added to the wells, and allowed to interact with the solid phase, which is the immobilized target molecule and scintillant coating in the wells. The assay can be measured immediately or allowed to reach equilibrium. Either way, when a radiolabel becomes sufficiently close to the scintillant coating, it produces a signal detectable by a device such as a TOPCOUNT NXT microplate scintillation counter (Packard BioScience Co., Meriden Conn.). If a radiolabeled expression product binds to the target molecule, the radiolabel remains in proximity to the scintillant long enough to produce a detectable signal.

[0835] In contrast, the labeled proteins that do not bind to the target molecule, or bind only briefly, will not remain near the scintillant long enough to produce a signal above background. Any time spent near the scintillant caused by random Brownian motion will also not result in a significant amount of signal. Likewise, residual unincorporated radiolabel used during the expression step may be present, but will not generate significant signal because it will be in solution rather than interacting with the target molecule. These non-binding interactions will therefore cause a certain level of background signal that can be mathematically removed. If too many signals are obtained, salt or other modifiers can be added directly to the assay plates until the desired specificity is obtained (Nichols et al., *Anal. Biochem.* 257:112-119, 1998).

[0836] XVI.E. Screening for Novel Antibiotics

[0837] As bacteria and other pathogens acquire resistance to known antibiotics, there is an ongoing interest in identifying novel antibiotics. See, e.g., Powell WA, Catranis CM, Maynard Calif. *Synthetic antimicrobial peptide design*. *Mol Plant Microbe Interact* 1995 September-October;8(5):792-4. Minicells can be used to assay, identify and purify novel antibiotics to eubacteria. By way of non-limiting example, a minicell that comprises a detectable compound can be contacted with a candidate antibiotic to see if the minicell is lysed by a candidate compound, which would release the detectable compound from the interior of the minicell into solution, this producing a signal that indicates that the candidate antibiotic is effective at lysing bacteria. In such assays, the detectable compound is such that it produces less or more of the same signal, or a different signal, inside the minicell as compared to in solution post-lysis. By way of non-limiting example, the minicell, could comprise a fluorescent compounds that, when contacted with a second fluorescent compound in solution, produces FRET.

[0838] XVI.F. Reverse Screening

[0839] In one version of minicell display, the invention provides methods for screening libraries of minicells in

which each minicell comprises an expression element that encodes a few, preferably one, membrane proteins in order to identify a membrane protein that interacts with a preselected compound. By way of non-limiting example, sequences encoding membrane proteins, fusion proteins, or cytoplasmic proteins are cloned into an expression vector, either by "shotgun" cloning or by directed cloning, e.g., by screening or selecting for cDNA clones, or by PCR amplification of DNA fragments, that encode a protein using one or more oligonucleotides encoding a highly conserved region of a protein family. For a non-limiting example of such techniques, see Krautwurst, D., et al. 1998. Identification of ligands for olfactory receptors by functional expression of a receptor library. *Cell* 95:917-926. By way of non-limiting example, a minicell expressing a receptor binds a preselected ligand, which may be a drug. Various assays for receptor binding, enzymatic activity, and channeling events are known in the art and may include detectable compounds; in the case of binding assays, competition assays may also be used (Masimirembwa, C. M., et al. 2001. *In vitro* high throughput screening of compounds for favorable metabolic properties in drug discovery. *Comb. Chem. High Throughput Screen.* 4:245-263; Mattheakis, L. C., and A. Saychenko. 2001. Assay technologies for screening ion channel targets. *Curr. Opin. Drug Discov. Devel.* 4:124-134; Numann, R., and P. A. Negulescu. 2001. High-throughput screening strategies for cardiac ion channels. *Trends Cardiovasc. Med.* 11:54-59; Le Poul, E., et al. 2002. Adaptation of aequorin functional assay to high throughput screening. *J. Biomol. Screen.* 7:57-65; and Graham, D. L., et al. 2001. Application of beta-galactosidase enzyme complementation technology as a high throughput screening format for antagonists of the epidermal growth factor receptor. *J. Biomol. Screen.* 6:401-411).

**[0840]** Once a minicell has been identified by an assay and isolated, DNA is prepared from the minicell. The cloned DNA present in the minicell encodes the receptor displayed by the minicell. Having been cloned, the receptor is used as a therapeutic target. For example, the receptor is produced via recombinant DNA expression and used in minicell-based or other assays to identify and characterize known and novel compounds that are ligands, antagonists and/or agonists of the cloned receptor. The ligands, antagonists and agonists may be used as lead compounds and/or drugs to treat diseases in which the receptor plays a role. In particular, when the preselected ligand is a drug, diseases for which that drug is therapeutic are expected to be treated using the novel ligands, antagonists and agonists, or drugs and prodrugs developed therefrom.

**[0841]** Preparations of minicells that express and secrete a soluble protein can be prepared in order to identify ligands, including but not limited to small molecules, that interact with the soluble protein. Soluble proteins include, but are not limited to, known secreted or proteolytically cleaved proteins and peptides, hormones and cytokines. In this format, minicells are placed in, or adhered to, the wells of a microtiter multiwell plate. A different compound or group of compounds is placed in each well, along with any reagents necessary to generate or quench a signal corresponding to a change in the soluble protein produced by the minicell. Such changes include, by way of non-limiting example, conformational changes in the protein that may occur as a result of binding of a ligand or otherwise. A well

that generates the appropriate signal contains a compound that causes a change in the soluble protein.

**[0842]** It is also possible to carry out procedures such as the one described in the immediately preceding paragraph "in reverse." In this format, a known ligand, which may be a drug, is used to identify soluble proteins that bind to the ligand/drug. Libraries of minicells wherein each minicell secretes a different soluble protein are prepared, and each type of minicell is placed into, or adhered to the wall of, a well of a microtiter plate, along with reagents for generating a signal when the ligand/drug binds to a soluble protein. Minicells that generate the appropriate signal comprise a cloned DNA that encodes a soluble protein that interacts with the known ligand/drug. Once cloned, the soluble protein is prepared and used as a therapeutic target in order to identify known or novel compounds that bind thereto. When the preselected ligand is a drug, diseases for which that drug is therapeutic are expected to be treated using the known and novel compounds so identified, or using drugs and prodrugs developed from such compounds.

**[0843]** Minicells expressing known membrane and soluble proteins can also be used to help characterize lead compounds and accelerate the generation of drugs therefrom. In particular, such studies may be used to identify potentially detrimental interactions that might occur upon *in vivo* administration, e.g., ADME/Tox screening (Ekins, S., et al. 2002. *In silico* ADME/Tox: the state of the art. *J. Mol. Graph. Model.* 20:305-309; and Li, A., et al. 2002. Early ADME/Tox studies and *in silico* screening. *Drug Discov. Today* 7:25-27).

**[0844]** By way of non-limiting example, a human receptor that is known to be important for the normal functioning of a cell may be expressed in minicells, and various chemical derivatives of a lead compound can be tested to ensure that they do not bind to the receptor, as such binding would be expected to have adverse effects *in vivo*. As another example, an enzyme that degrades a drug, such as cytochrome P450, is expressed in minicells and used to examine the susceptibility of a candidate drug to such degradation. The cytochrome P450 family of enzymes is primarily responsible for the metabolism of xenobiotics such as drugs, carcinogens and environmental chemicals, as well as several classes of endobiotics such as steroids and prostaglandins. Exemplary P450 cytochromes involved in drug degradation include, but are not limited to, CYP2D6 (cytochrome P4502D6, also known as debrisoquine hydroxylase), CYP1A1, CYP1A2, CYP2A6, CYP2B6, CYP2C9, CYP2C18, CYP2C19, CYP2D6, CYP2E1, CYP3A4 and CYP3A5.

**[0845]** XVI.G. Molecular Variants

**[0846]** In one aspect of the invention, minicells are used in methods of screening to identify agents that improve, enhance, or decrease the interaction of a protein with another compound. These methods include, by way of non-limiting example, modification of protein targets through directed or random mutagenic approaches to identify critical interactions between a wild-type protein target and a specific drug molecule. Information obtained from studies of mutant proteins is used to specifically produce or modify a therapeutic agent to interact more specifically and/or effectively with the wild-type protein target, thus increasing the therapeutic efficacy of the parental drug and/or decreasing non-

specific, potentially deleterious interactions. See, for example, Lietha, D., et al. 2001. Crystal structures of NK1-heparin complexes reveal the basis for NK1 activity and enable engineering of potent agonists of the MET receptor. *EMBO J.* 20:5543-5555; and Chen, Y. Z., et al. Can an optimization/scoring procedure in ligand-protein docking be employed to probe drug-resistant mutations in proteins? *J. Mol. Graph. Model.* 19:560-570; Zhao, H. and F. H. Arnold. Combinatorial protein design: Strategies for screening protein libraries. *Current Opinion in Structural Biology* 7:480-485 (1997); and Carrupt P A, el Tayar N, Karlen A, Testa B. Molecular electrostatic potentials for characterizing drug-biosystem interactions. *Methods Enzymol.* 1991;203:638-77. Martin Y C. Computer-assisted rational drug design. *Methods Enzymol.* 1991;203:587-613.

**[0847]** By way of non-limiting example, information obtained using the methods of the invention may be in conjunction with x-ray crystallographic structural determinations to characterize receptor:ligand interactions (Muller, G. 2000. Towards 3D structures of G protein-coupled receptors: a multidisciplinary approach. *Curr. Med. Chem.* 7:861-888). By way of non-limiting example, minicells may be used to display the family of molecular variants to characterize the specific mutagenic changes on the functional properties of the protein.

**[0848]** Studies of variant proteins can also be used to modify drugs to fit natural variants of proteins, especially those associated with pathogens. Pathogens such as viruses, including retroviruses such as HIV, may acquire mutations that change a site where a drug acts, thereby rendering the pathogen immune to the drug. Studies of variant proteins can be used to quickly produce derivatives of a drug that are active against a variant protein. See, for example, Varghese J N, Smith P W, Sollis S L, Blick T J, Sahasrabudhe A, McKimm-Breschkin J L, Colman P M. Drug design against a shifting target: a structural basis for resistance to inhibitors in a variant of influenza virus neuraminidase. *Structure Jun.* 15, 1998;6(6):735-46; and Baldwin E T, Bhat T N, Liu B, Pattabiraman N, Erickson J W. Structural basis of drug resistance for the V82A mutant of HIV-1 proteinase. *78: Nat Struct Biol* 1995 March;2(3):244-9.

**[0849]** XVI.H. Directed Evolution

**[0850]** The minicells and methods described herein can be used in directed evolution. Unlike natural variation, directed evolution generates new protein variants in vitro (see, e.g., Arnold, F. H. and A. A. Violkov. Directed Evolution of Biocatalysts. *Curr Op Chem Biol* 1999. 3:54-59). Amino acid substitutions can be introduced into a protein of interest by mutating the gene encoding the protein. Mutations are introduced by, e.g., replicating DNA in mutator strains, by chemical mutagenesis or radiation-induced mutagenesis (Drake, J. W., *The Molecular Basis of Mutation*, Holden-Day, San Francisco, 1970). Other methods include error-prone PCR and "domain shuffling" (Moore, G. L. and C. D. Maranas. Modeling DNA Mutation and Recombination for Directed Evolution Experiments. *J. Ther. Biol.* 2000. 205:483-503). In the latter method, different regions of members of the same gene family are recombined so that the inherent variability of members of the family is used to produce novel "isoforms" of genes.

**[0851]** A group of variants is screened to select for those variants which have the desired activity. The activity of the

initial variants that are so isolated may be inadequate for a given application, but the process can be repeated using these initial members to generate a second group of variants, or reiterated as many times as is necessary to produce one or more variants having the desired activity or characteristics.

**[0852]** XVI.I. Isolation and Characterization of Components of Signal Transduction Pathways

**[0853]** In one version of minicell display, the invention provides methods for screening libraries of minicells, in which each minicell comprises a preselected component of a signal transduction pathway, in order to identify soluble and membrane proteins that interact with the preselected component. By way of non-limiting example, a plurality of minicells, each of which displays the same G-protein-coupled receptor (GPCR), is used to prepare a minicell library in which a different G-protein encoding sequence is present and expressed in each minicell. Minicells comprising a G-protein that interacts with the GPCR are identified, e.g., via transactivation assays described in Example 18. Once a minicell has been identified by an assay and isolated, DNA is prepared from the minicell. The cloned DNA present in the minicell encodes a G-protein that interacts with the GPCR of the displayed by the minicells of the library. Having been cloned, the G-protein is used as a therapeutic target that can be used in screening assays to identify novel lead compounds and drugs that interfere or alter the activity of the GPCR. In particular, when the GPCR of the minicell library is known to be a therapeutic target for a specific disease, it is expected that compounds that interfere or alter the activity of a G-protein that interacts with the GPCR will be or lead to therapeutics for that specific disease.

**[0854]** In addition to G-protein signal transduction pathways, other non-limiting examples of signal transduction pathways include the MAPK pathway, the SAPK pathway, the p38 pathway and/or the ceramide-mediated stress response pathway. See Zhang, W., and L. E. Samelson. 2000. The role of membrane-associated adaptors in T cell receptor signalling. *Semin. Immunol.* 12:35-41; Liebmann, C. 2001. Regulation of MAP kinase activity by peptide receptor signalling pathway: paradigms of multiplicity. *Cell Signal.* 13:777-785; Lee, Jr., J. T., and J. A. McCubrey. 2002. The Raf/MEK/ERK signal transduction cascade as a target for chemotherapeutic intervention in leukemia. *Leukemia.* 16:486-507; Tibbles, L. A., and J. R. Woodgett. 1999. The stress-activated protein kinase pathways. *Cell Mol. Life Sci.* 55:1230-1254; Rao, K. M. 2001. MAP kinase activation of macrophages. *J. Leukoc. Biol.* 69:3-10; Pelech, S. L., and D. L. Charest. 1995. MAP kinase-dependent pathways in cell cycle control. *Prog. Cell Cycle Res.* 1:33-52; Lee, S. H., et al. 2001. BetaPix-enhanced p38 activation by Cdc42/Rac/PAK/MKK3/6-mediated pathway. Implication in the regulation of membrane ruffling. *J. Biol. Chem.* 276:25066-25072; Ono, K., et al. 2000. The p38 signal transduction pathway Activation and function. *Cellular Signalling* 12:1-13; You, A. 2001. Differentiation, apoptosis, and function of human immature and mature myeloid cells: intracellular signaling mechanism. *Int. J. Hematol.* 73:438-452; Johnson, D. I. 1999. Cdc42: An essential Rho-type GTPase controlling eukaryotic cell polarity. *Microbiol. Mol. Biol. Rev.* 63:54-105; Williams, J. A. 2001. Intracellular signaling mechanisms activated by cholecystokinin-regulating synthesis and secretion of digestive enzymes in pancreatic acinar cells. *Annu. Rev. Physiol.* 63:77-97; Mathias, S., et al.

1998. Signal transduction of stress via ceramide. *Biochem J.* 335:465-480; and Hannun, Y. A., et al. 2000. Ceramide in the eukaryotic stress response. *Trends Cell Biol.* 10:73-80.

[0855] XVII. Determining the Structures of Membrane Proteins

[0856] Three-dimensional (3D) structures of proteins may be used for drug discovery. However, GPCRs and other membrane proteins present challenging problems for 3D structure determination. Muller, Towards 3D structures of G protein-coupled receptors: a multidisciplinary approach. (Review), *Curr Med Chem* 2000 pp.861-88; Levy et al., Two-dimensional crystallization on lipid layer: A successful approach for membrane proteins, *J Struct Biol* 1999 127, 44-52. Although the three-dimensional structures of hundreds of different folds of globular proteins have been determined, fewer than 20 different integral membrane protein structures have been determined. There are many reasons for this. Extracting membrane proteins from the membrane can easily disrupt their native structure, and membrane proteins are notoriously difficult to crystallize.

[0857] Some membrane proteins readily form two-dimensional crystals in membranes and can be used for structure determination using electron diffraction spectroscopy (ED) instead of x-ray crystallography. This is the technique that was used to determine the structure of bacteriorhodopsin (see below).

[0858] Nuclear magnetic resonance (NMR) is an alternative method for determining membrane protein structure, but most membrane proteins are too large for high-resolution NMR at the present state of the art. Furthermore, membrane proteins require special conditions for NMR, e.g. deuterated lipids must be used to avoid confusing the signal of the protein protons with the noise of membrane lipid protons.

[0859] Membrane protein for which structures have been determined include photosynthetic reaction center, porin, porin OmpF, plant light-harvesting complex (chlorophyll a-b binding protein), bacterial light-harvesting complex, cytochrome c oxidase, glycophorin A, the Sec A translocation ATPase of *Bacillus subtilis*, and a bacterial potassium channel. For details, see: Weinkauff et al., (2001): Conformational stabilization and crystallization of the Sec A translocation ATPase from *Bacillus subtilis*. *Acta Crystallogr D Biol Crystallogr* 57:559-565; Cowan et al., (1992): Crystal structures explain functional properties of two *E. coli* porins. *Nature* 358:727-33; Deisenhofer et al., (1984): X-ray structure analysis of a membrane protein complex. Electron density map at 3 Å resolution and a model of the chromophores of the photosynthetic reaction center from *Rhodospseudomonas viridis*. *J Mol Biol* 180:385-98; Deisenhofer et al., (1985): Structure of the protein subunits in the photosynthetic reaction centre of *Rhodospseudomonas viridis* at 3 Å resolution. *Nature* 318:618; Doyle et al., (1998): The structure of the potassium channel: molecular basis of K<sup>+</sup> conduction and selectivity. *Science* 280:69-77; Henderson et al., (1990): Model for the structure of bacteriorhodopsin based on high-resolution electron cryo-microscopy. *J Mol Biol* 213:899-929; Iwata et al., (1998): Complete structure of the 11-subunit bovine mitochondrial cytochrome bc<sub>1</sub> complex. *Science* 281:64-71; Koepke et al., (1996): The crystal structure of the light-harvesting complex II (B800-850) from *Rhodospirillum rubrum*. *Structure* 4:581-97; Kuhlbrandt et al., (1994): Atomic model of

plant light-harvesting complex by electron crystallography. *Nature* 367:614-21; Lemmon et al., (1992): Sequence specificity in the dimerization of transmembrane alpha-helices. *Biochemistry* 31:12719-25; MacKenzie et al., (1997): A transmembrane helix dimer: structure and implications. *Science* 276:131-3; McDermott et al., (1995): Crystal structure of an integral membrane light-harvesting complex from photosynthetic bacteria, *Nature* 374:517-21; Michel (1982): Three-dimensional crystals of a membrane protein complex. The photosynthetic reaction centre from *Rhodospseudomonas viridis*. *J Mol Biol* 158:567-72; Tsukihara et al., (1996): The whole structure of the 13-subunit oxidized cytochrome c oxidase at 2.8 Å. *Science* 272:1136-44; and Weiss et al., (1991): The structure of porin from *Rhodobacter capsulatus* at 1.8 Å resolution. *FEBS Lett* 280:379-82. Table 5, which is based upon Preusch et al. (1998) as revised by White & Wimley (1999), lists membrane proteins whose crystallographic structures have been determined.

TABLE 5

Structural Data Regarding Membrane Proteins	
PROTEIN	REFERENCES
<b>MONOTOPIC MEMBRANE PROTEINS</b>	
Portaglandin H2 synthase-1. Sheep. 3.5Å	Picot et al. (1994)
Cyclooxygenase-2. <i>Mus Musculus</i> . 3.0Å	Kurumbail et al. (1996)
Squalene-hopene cyclase. <i>Alicyclobacillus acidocaldarius</i> . 2.0Å	Wendt et al. (1999)
<b>TRANSMEMBRANE PROTEINS</b>	
Bacterial Rhodopsins ( <i>Halobacterium salinarium</i> )	
Bacteriorhodopsin (BR)	
2D xtals. EM. 3.5Å	Grigorieff et al. (1996)
2D xtals. EM. 3.0Å	Kimura et al. (1997)
3D xtals. X-ray. 2.5Å	Pebay-Peyroula et al. (1997)
3D xtals. X-ray. 1.9Å	Belhrhali et al. (1999)
3D xtals. X-ray 2.1Å K intermediate	Edman et al. (1999)
3D xtals. X-ray. 2.3Å	Luecke et al. (1998)
3D xtals. X-ray. 1.55Å	Luecke et al. (1999)
3D xtals. X-ray. D96N mutant (BR) 1.80Å	Luecke et al. (1999)
3D xtals. X-ray. D96N mutant (M) 2.00Å	
3D xtals. X-ray. 2.9Å	Essen et al. (1998)
Halorhodopsin (HR)	
3D xtals. X-ray. 1.8Å	Kolbe et al. (2000)
<b>G PROTEIN-COUPLED RECEPTORS</b>	
Rhodopsin. Bovine Rod Outer Segment. 2.8Å	Palczewski et al. (2000)
<b>Photosynthetic Reaction Centers</b>	
<i>Rhodospseudomonas viridis</i> . 2.3Å	Deisenhofer et al. (1985)
<i>Rhodobacter sphaeroides</i> . 3.0Å	Yeates et al. (1987)
<i>Rhodobacter sphaeroides</i> . 3.1Å	Chang et al. (1991)

TABLE 5-continued

<u>Structural Data Regarding Membrane Proteins</u>	
PROTEIN	REFERENCES
<u>Light Harvesting Complexes</u>	
<i>Rhodospseudomonas acidophyla</i> . 2.5Å	McDermott et al. (1995)
<i>Rhodospirillum molitischianum</i> . 2.4Åk	Koepke et al. (1996)
<u>Photosystems</u>	
Photosystem I. <i>Synechococcus elongates</i> 4.0Å	Schubert et al. (1997)
Photosystem II. <i>Synechococcus elongates</i> 3.8Å	Zouni et al. (2001)
<u>Beta-Barrel Membrane Proteins-Multimeric (Porins and Relatives)</u>	
Porin. <i>Rhodobacter capsulatus</i> . 1.8Å	Weiss & Schulz (1992)
Porin. <i>Rhodopseudomonas blastica</i> 1.96Å	Kreutsch et al. (1994)
OmpF. <i>E. coli</i> . 2.4Å	Cowan et al. (1992)
PhoE. <i>E. coli</i> . 3.0Å	Cowan et al. (1992)
Maltoporin. <i>Salmonella typhimurium</i> . 2.4Å	Meyer et al. (1997)
Maltoporin. <i>E. coli</i> 3.1Å	Schirmer et al. (1995)
<u>Beta-Barrel Membrane Proteins-Monomeric/Dimeric</u>	
ToIC outer membrane protein. <i>E. coli</i> 2.1Å Protein is a trimer, each contributing 4 strands to a single barrel.	Koronakis et al. (2000)
OmpA. <i>E. coli</i> . 2.5Å	Pautsch & Schulz (1998)
OmpA. <i>E. coli</i> . By NMR, in DPC micelles	Arora et al. (2001)
OmpX. <i>E. coli</i> . 1.9Å	Vogt & Schulz (1990)
OMPLA (outer membrane phospholipase A) <i>E. coli</i> . 2.1Å. monomer (1QD5) and dimer (1QD6)	Snijder et al. (1999)
FhuA. <i>E. coli</i> . 2.5Å	Ferguson et al. (1998); Lambert et al., 1999
FhuA + ferrichrome-iron. <i>E. coli</i> . 2.7Å	Buchanan et al. (1999)
FepA. <i>E. coli</i> . 2.4Å	Ferguson et al. (1999)
Glycophorin A. humanm.	MacKenzie et al. (1997)
<u>Non-constitutive Toxins, etc.</u>	
Alpha-hemolysin. <i>Staphylococcus aureus</i> . 1.9Å	Song et al. (1996)
LukF. <i>Staphylococcus aureus</i> . 1.9Å	Olson et al. (1999)
<u>Ion Channels</u>	
KcsA Potassium, H <sup>+</sup> gated. <i>Streptomyces lividans</i> . 3.2Å	Doyle et al. (1998)
MscL Mechanosensitive. <i>Mycobacterium tuberculosis</i> . 3.5Å	Chang et al. (1998)
<u>Other Channels</u>	
AQP1 - aquaporin water channel. Red blood cell. Electron crystallography in membrane plane. 3.8Å	Murata et al. (2000)
AQP1 - In vitreous ice by electron microscopy. 3.7Å	Ren et al. (2000)
GipF - glycerol facilitator channel. <i>E. coli</i> . 2.2Å	Fu et al. (2000)
<u>P-type ATPase</u>	
Calcium ATPase. <i>Sarcoplasmic reticulum</i> . Rabbit. 2.6Å	Toyoshima et al. (2000)

TABLE 5-continued

<u>Structural Data Regarding Membrane Proteins</u>	
PROTEIN	REFERENCES
<u>Respiratory Proteins</u>	
Fumarate Reductase Complex. <i>Escherichia coli</i> . 3.3Å	Iverson et al. (1999)
Fumarate Reductase Complex. <i>Wolinella succinogenes</i> . 2.2Å	Lancaster et al. (1999)
ATP synthase (F <sub>1c10</sub> ). <i>S. cerevisiae</i> . 3.9Å. X-ray structure is a C alpha model derived from composite of 1BMF, 1A91 & 1AQT	Stock et al. (1999)
<u>Cytochrome C Oxidases</u>	
aa <sub>3</sub> bovine heart mitochondria. 2.8Å	Tsukihara et al. (1996)
aa <sub>3</sub> <i>Paracoccus denitrificans</i> . 2.8Å	Iwata et al. (1995)
ba <sub>3</sub> from <i>T. thermophilus</i> . 2.4Å	Soulimane et al. (2000)
<u>Cytochrome b<sub>c1</sub> Complexes</u>	
Bovine Heart Mitochondria (5 subunits). 2.9Å	Xia et al. (1997)
Chicken Heart Mitochondria. 3.16Å	Zhang et al. (1998)
Bovine Heart Mitochondria (11 subunits). 2.8-3.0Å.	Iwata et al. (1998)
<i>S. cerevisiae</i> (yeast, 9 subunits). 2.3Å	Hunte et al. (2000)

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- [0914] XVIII. Biosensors and Environmental Applications
- [0915] XVIII.A. Minicell-Based Biosensors
- [0916] The present invention is directed to a device that comprises a sensor adapted to detect one or more specific health and/or nutrition markers in a subject or in the environment. The device may also signal the caretaker, the subject, or an actuator of the occurrence. The sensor comprises a biosensor. As used herein, the term "biosensor" is defined as a component comprising one or more binding moieties being adapted to detect a ligand found in one or more target pathogenic microorganisms or related biomolecules.
- [0917] Generally, biosensors function by providing a means of specifically binding, and therefore detecting, a target biologically active analyte. In this way, the biosensor is highly selective, even when presented with a mixture of many chemical and biological entities. Often the target biological analyte is a minor component of a complex mixture comprising a multiplicity of biological and other components. Thus, in many biosensor applications, detection of target analytes occurs in the parts-per-billion, parts-per-trillion, or even lower ranges levels.
- [0918] XVIII.A.1. Minicell-Based Biosensor Design
- [0919] The biosensor of the present invention may comprise a bio-recognition element, or molecular recognition element, that provides the highly specific binding or detection selectivity for a particular analyte. In a biosensor of the invention, the bio-recognition element, or system, is a minicell displaying an enzyme or sequence of enzymes; an antibody or antibody derivative; a membrane receptor protein; or the like, and generally functions to interact specifically with a target biological analyte. The bio-recognition element is responsible for the selective recognition of the analyte and the physico-chemical signal that provides the basis for the output signal. The expressed protein or molecule does not need to be a naturally occurring membrane bound protein but could be a soluble protein or small molecule tethered to the minicell by, for example, a transmembrane domain of another protein such as the EGFR or ToxR.
- [0920] Biosensors may include biocatalytic biosensors, and bioaffinity biosensors. In biocatalytic biosensor embodiments, the bio-recognition element minicell is "biocatalytic," e.g., displays an enzyme. In biocatalytic biosensors, the selective binding sites "turn over" (i.e., can be used again during the detection process), resulting in a significant amplification of the input signal. Biocatalytic sensors such as these are generally useful for real-time, continuous sensing.
- [0921] Bioaffinity sensors are generally applicable to bacteria, viruses, toxins and other undesirable compounds and include chemoreceptor-based biosensors and/or immunological sensors (i.e., immunosensors). Chemoreceptors are complex biomolecular macroassemblies responsible, in part, for a viable organism's ability to sense chemicals in its environment with high selectivity. Chemoreceptor-based biosensors comprise one or more natural or synthetic chemoreceptors associated with a means to provide a signal (visual, electrical, etc.) of the presence or concentration of a target biological analyte. In certain embodiments, the chemoreceptor may be associated with an electrode (i.e., an electrical transducer) so as to provide a detectable electrical signal. In the biosensors of the invention, minicells displaying a receptor are used in place of chemoreceptors. The minicell has many desired features of a viable cell, and performs similar functions, but is more durable.
- [0922] On the other hand, the bio-recognition elements of immunosensors are generally antibodies or antibody derivatives. In any case, bioaffinity biosensors are generally irreversible because the receptor sites of the biosensor become saturated when exposed to the target biological analyte. In a biosensor of the invention, an immunosensor may be a minicell displaying an antibody or antibody fragment.
- [0923] Biocatalytic and bioaffinity biosensor systems are described in more detail in *Journal of Chromatography*, 510 (1990) 347-354 and in the *Kirk-Othmer Encyclopedia of Chemical Technology*, 4.sup.th ed. (1992), John Wiley & Sons, NY, the disclosure of which is incorporated by reference herein.
- [0924] The biosensors of the present invention may detect biologically active analytes related to impending (i.e., future presentation of symptoms is likely) or current human systemic disease states, including, but not limited to, pathogenic bacteria, parasites (e.g., any stage of the life cycle, including eggs or portions thereof, cysts, or mature organisms), viruses, fungi such as *Candida albicans*, antibodies to pathogens, and/or microbially produced toxins. Additionally, the biosensor may target biologically active analytes

related to impending or current localized health issues, such as stress proteins (e.g., cytokines) and interleukin 1-alpha that may precede the clinical presentation of skin irritation or inflammation. In preferred embodiments, the biosensor functions as a proactive sensor, detecting and signaling the subject, a caretaker or medical personnel of the impending condition prior to the presentation of clinical symptoms. This allows time to administer prophylactic or remedial treatments to the subject which can significantly reduce, if not prevent, the severity and duration of the symptoms. Further, the sensor, by detecting the presence of a target biological analyte in a sample from the subject, may detect residual contamination on a surface, such as skin or environmental surface, in contact with the biosensor, and provide and appropriate signal.

[0925] The physico-chemical signal generated by the bio-recognition element or elements may be communicated visually to the caretaker or medical personnel (i.e., via a color change visible to the human eye). Other embodiments may produce optical signals, which may require other instrumentation to enhance the signal. These include fluorescence, bioluminescence, total internal reflectance resonance, surface plasmon resonance, Raman methods and other laser-based methods, such as LED or laser diode sensors. For example, exemplary surface plasmon resonance biosensors are available as IBIS I and IBIS II from XanTec Analysensysteme of Muenster, Germany, which may comprise bioconjugate surfaces as bio-recognition elements. Alternatively, the signal may be processed via an associated transducer which, for example, may produce an electrical signal (e.g., current, potential, inductance, or impedance) that may be displayed (e.g., on a readout such as an LED or LCD display) or which triggers an audible or tactile (e.g., vibration) signal or which may trigger an actuator, as described herein. The signal may be qualitative (e.g., indicating the presence of the target biological analyte) or quantitative (i.e., a measurement of the amount or concentration of the target biological analyte). In such embodiments, the transducer may optionally produce an optical, thermal or acoustic signal.

[0926] In any case, the signal may also be durable (i.e., stable and readable over a length of time typically at least of the same magnitude as the usage life of the device) or transient (i.e., registering a real-time measurement). Additionally, the signal may be transmitted to a remote indicator site (e.g., via a wire, or transmitter, such as an infrared or rf transmitter) including other locations within or on the device or remote devices. Further, the sensor, or any of its components, may be adapted to detect and/or signal only concentrations of the target biological analyte above a predefined threshold level (e.g., in cases wherein the target biological analyte is normally present in the bodily waste or when the concentration of the analyte is below a known "danger" level).

[0927] The target analytes that the biosensors of the present invention are adapted to detect may also be viruses. These may include diarrhea-inducing viruses such as rotavirus, or other viruses such as rhinovirus and human immunodeficiency virus (HIV). An exemplary biosensor adapted to detect HIV is described in U.S. Pat. Nos. 5,830,341 and 5,795,453, referenced above. The disclosure of each of these patents is incorporated by reference herein. Biosensors are adapted to use in different tissues; see, e.g., U.S. Pat. No. 6,342,037; Roe et al. Jan. 29, 2002; Device having fecal

component sensor; and using different binding molecules, see, e.g., U.S. Pat. No. 6,329,160; Schneider et al. Dec. 11, 2001; Biosensors.

[0928] When minicells are incorporated into a biosensor, they may be immobilized in the biosensor by techniques known in the art such as entrapment, adsorption, crosslinking, encapsulation, covalent attachment, any combination thereof, or the like. Further, the immobilization can be carried out on many different substrates such as known the art. In certain preferred embodiments, the immobilization substrate may be selected from the group of polymer-based materials, hydrogels, tissues, nonwoven materials or woven materials.

[0929] In certain embodiments, biosensor embodiments, may comprise, be disposed on, or be operatively associated with a microchip, such as a silicon chip, MEMs (i.e., micro electromechanical system) device, or an integrated circuit. Microchip-based biosensors may be known as "biochips". Regardless of the type of sensor, the microchip may comprise a multiplicity of sensor components having similar or different sensitivities, kinetics, and/or target analytes (i.e., markers) in an array adapted to detect differing levels or combinations of the analyte(s). Further, each sensor in such an array may provide a different type of signal, including those types disclosed herein, and may be associated with different actuators and/or controllers. Also, each sensor in an array may operate independently or in association with (e.g., in parallel, combination, or series) any number of other sensors in the array.

[0930] A minicell of a biosensor of the invention may comprise a detectable compound that produces a signal once ligands have bound to the minicell. By way of non-limiting example, a minicell may display a receptor for a ligand and contain a fluorescent compound. The binding and internalization of the ligand into the minicell results in FRET, shifting the wavelength of the signal. See, by way of non-limiting example, Billinton et al., Development of a green fluorescent protein reporter for a yeast genotoxicity biosensor, *Biosensors & Bioelectronics* 13:831-838, 1998. A biosensor according to the invention may use microbalance sensor systems (Hengerer et al., Determination of phage antibody affinities to antigen by a microbalance sensor system, *BioTechniques* 26:956-964, 1999).

[0931] XVIII.A.2. Surface Plasmon Resonance

[0932]  $K_d$  is measured using surface plasmon resonance on a chip, for example, with a BIAcore® chip coated with immobilized binding components, or similar systems such as the IAsys from Thermo Labsystems, Affinity Sensors Division (Cambridge, U.K.) or the BIOS-1 system from Artificial Sensing, Inc. (Zurich, Switzerland). See Fitzgerald, Coupling optical biosensor technology with micro-preparative HPLC: Part 1, *Am Biotech Lab* November 2000, p.10 and 12; Fitzgerald, Coupling optical biosensor technology with micro-preparative HPLC: Part 2, *Am Biotech Lab* Feb. 14, 16 and 18, 2001; and Leatherbarrow et al., Analysis of molecular recognition using optical sensors, *Current Opinion in Chem Biol* 3:544-547, 1999).

[0933] Surface plasmon resonance is used to characterize the microscopic association and dissociation constants of reaction between an antibody or antibody fragment and its ligand. Such methods are generally described in the follow-



ing references that are incorporated herein by reference. (Vely F. et al., BIAcore analysis to test phosphopeptide-SH2 domain interactions, *Methods in Molecular Biology*. 121:313-21, 2000; Liparoto et al., Biosensor analysis of the interleukin-2 receptor complex, *Journal of Molecular Recognition*. 12:316-21, 1999; Lipschultz et al., Experimental design for analysis of complex kinetics using surface plasmon resonance, *Methods*. 20(3):310-8, 2000; Malmqvist., BIAcore: an affinity biosensor system for characterization of biomolecular interactions, *Biochemical Society Transactions* 27:335-40, 1999; Alftan, Surface plasmon resonance biosensors as a tool in antibody engineering, *Biosensors & Bioelectronics*. 13:653-63, 1998; Fivash et al., BIAcore for macromolecular interaction, *Current Opinion in Biotechnology*. 9:97-101, 1998; Price et al.; Summary report on the ISOBM TD-4 Workshop: analysis of 56 monoclonal antibodies against the MUC1 mucin. *Tumour Biology* 19 Suppl 1:1-20, 1998; Malmqvist et al, Biomolecular interaction analysis: affinity biosensor technologies for functional analysis of proteins, *Current Opinion in Chemical Biology*. 1:378-83, 1997; O'Shannessy et al., Interpretation of deviations from pseudo-first-order kinetic behavior in the characterization of ligand binding by biosensor technology, *Analytical Biochemistry*. 236:275-83, 1996; Malmberg et al., BIAcore as a tool in antibody engineering, *Journal of Immunological Methods*. 183:7-13, 1995; Van Regenmortel, Use of biosensors to characterize recombinant proteins, *Developments in Biological Standardization*. 83:143-51, 1994; O'Shannessy, Determination of kinetic rate and equilibrium binding constants for macromolecular interactions: a critique of the surface plasmon resonance literature, *Current Opinions in Biotechnology*. 5:65-71, 1994).

[0934] BIAcore® uses the optical properties of surface plasmon resonance (SPR) to detect alterations in protein concentration bound within to a dextran matrix lying on the surface of a gold/glass sensor chip interface, a dextran biosensor matrix. In brief, proteins are covalently bound to the dextran matrix at a known concentration and a ligand for the protein (e.g., antibody) is injected through the dextran matrix. Near infra red light, directed onto the opposite side of the sensor chip surface is reflected and also induces an evanescent wave in the gold film, which in turn, causes an intensity dip in the reflected light at a particular angle known as the resonance angle. If the refractive index of the sensor chip surface is altered (e.g., by ligand binding to the bound protein) a shift occurs in the resonance angle. This angle shift can be measured and is expressed as resonance units (RUs) such that 1000 RUs is equivalent to a change in surface protein concentration of 1 ng/mm<sup>2</sup>. These changes are displayed with respect to time along the y-axis of a sensorgram, which depicts the association and dissociation of any biological reaction.

[0935] Additional details may be found in Jonsson et al., Introducing a biosensor based technology for real-time biospecific interaction analysis, (1993) *Ann. Biol. Clin.* 51:19-26; Jonsson et al., Real-time biospecific interaction analysis using surface plasmon resonance and a sensor chip technology, (1991) *Biotechniques* 11:620-627; Jonsson et al., Comparison of methods for immobilization to carboxymethyl dextran sensor surfaces by analysis of the specific activity of monoclonal antibodies, (1995) *J. Mol. Recognit.* 8:125-131; and Jonsson, Immobilization of proteins to a carboxymethyl dextran-modified gold surface for biospecific interaction analysis in surface plasmon reso-

nance sensors (1991) *Anal. Biochem.* 198:268-277, Karlsson et. al., Kinetic analysis of monoclonal antibody-antigen interactions with a new biosensor based analytical system *J. Immunol. Meth.*, 145, 229, 1991; Weinberger et al., Recent trends in protein biochip technology, *Pharmacogenomics* 2000 November; 1(4):395-416; Lipschultz et al., Experimental design for analysis of complex kinetics using surface plasmon resonance, *Methods* 2000 March;20(3):310-8.

#### [0936] XVIII.B. Toxicological Sampling

[0937] Minicells are ideally suited for in vitro diagnostic toxicological applications in which toxins, poisons, infectious agents or pathogens, heavy metals, pollutants, caustic agents, allergens, organic molecules, radionuclides, or other environmental contaminants present either in air, water, soil samples and/or fluid and/or tissue samples of organisms can be assessed. An embodiment of this invention, minicells expressing proteins or other molecules could be used in variety of diagnostic detection platforms, including microwell formats, lateral flow devices, molecular switches, biosensors, badges and other sensing devices. Without being limited to the following examples, such devices could be used for early warning of chemical and/or bioweapon attack, illegal drug detection, explosives detection, biohazard detection, pollution assessment, pesticide contamination, allergen detection and detection of toxic or hazardous gasses. In a related application, minicells could be used to eliminate, modify or inactivate the agents.

[0938] In one non-limiting example of protein expression on minicells for toxicological detection, olfactory receptors could be expressed by minicells. The olfactory system possesses the ability to recognize and differentiate between a wide range of odorants based on odor molecules interacting with specific receptor proteins in the ciliary membrane of olfactory neurons (Lancet, D., 1986. Vertebrate olfactory reception. *Ann. Rev. Neurosci.* 9:329-355; Shepherd, G. M., 1994. Discrimination of molecular signals by the olfactory receptor neuron. *Neuron* 13:771-790). These receptors were found to be 7-transmembrane-domain members of the G protein-coupled receptor family (Buck, L. and R. Alex. 1991. A novel multigene family may encode odorant receptors: A molecular basis for odor recognition. *Cell* 65:175-187). Using a murine receptor library, olfactory receptors were functionally expressed in HEK-293 cells (Krautwurst, D., et al., 1998. Identification of ligands for olfactory receptors by functional expression of a receptor library. *Cell*. 95:917-926). By coexpressing the cloned receptors with G 15,16 subunits, the modified receptor system upon activation leads to an increase in intracellular Ca<sup>2+</sup>. Calcium levels were measured employing the dye FURA-2 and ratiofluorimetric imaging. This system demonstrated ligand specificity and structure-function relationships for identified olfactory receptors. Employing similar techniques, OR17-40, a human olfactory receptor protein, was expressed in human embryonic kidney 293 cells and *Xenopus Laevis* oocytes (Wetzal, H., et al. 1999. Specificity and sensitivity of a human olfactory receptor functionally expressed in Human Embryonic Kidney 293 Cells and *Xenopus Laevis* Oocytes. *J. Neurosciences*. 19:7426-7433). The receptor was functionally expressed in a manner designed to assess the, specificity of its binding to the ligand, helional.

[0939] In one non-limiting example of target protein identification, primers from homologous areas in transmembrane

II and transmembrane VII of olfactory GPCRs will be used to identify unique receptor sequences. These sequences are inserted into expression vectors. Minicell producing bacteria are transformed with these vectors and cultured. Minicells are isolated from the culture as previously described and subsequently induced. Using HTS previously described, the functional receptor/minicells which generate signal for binding of an odiferous toxin to the receptor are identified. Large scale production of the minicells is carried out and the minicells are covalently coupled to the surface of a microarray chip. The chip is supported in an air sampler, which feeds atmosphere over surface of the chip on a continuous basis. If the toxic agent is present in the air, the binding to the receptor activates a series of events ending in the generation of a signal identifying the presence of the agent in the air.

[0940] By way of non-limiting example, standard molecular biological techniques can be used as follows: cDNA for GFP is ligated to the 3' end of cDNA sequence for the receptor described above. The resulting sequence is inserted into an expression vector. Minicell producing bacteria are transformed with these vectors and cultured. Minicells are isolated from the culture as previously described and subsequently induced. The minicells now contain the receptor to the ligand on the surface of the minicell with a GFP tag on the C-terminus of the protein in the cytosol. These minicells are packed into filters. Air is passed through the filter. If the ligand is present, it will bind to the receptor. The filter packing is suspended on applied to a diagnostic device. Antibody to the ligand/receptor binding site complex is fixed on the capture zone. When the sample is applied to the device, the receptor/ligand complex is captured. The capture zone is screened for signal resulting from the presence of GFP. This can be extrapolated to have multiple unique receptor/minicell moieties in the same sampling device. Each receptor would have a unique fluorescing protein tag such that different emissions identify specific agents in the air.

[0941] Other methods for quantification associated take advantage of the composition of the minicell. Loading of the minicell by transiently permeabilizing the membrane to allow for migration of molecules into to the cytosol. These molecules include but are not limited to radiolabeled molecules (i.e., nucleotides), stains or dyes (DAPI or other DNA staining, heavy metals, fluorophores. The molecules could also be synthesized within the minicell (i.e. GFP). The association of a specific ligand with the minicell could cause a redox shift that induce a color change in the solution or could shift the energy potential in the reaction are generating an electrical current. Each of this examples are associated with well know methods for measuring each of the resulting changes. These include but are not limited to radioactivity or fluorescence generated or the color shift by spectrophotometry.

[0942] A multigene family of gustatory G protein-coupled receptors expressed in the lingual epithelia has been identified with structural similarities to olfactory receptors (Abe, K., et al. 1993. Multiple genes for G protein-coupled receptors and their expression in lingual epithelia. FEBS. 316:253-256; Abe, K., et al. 1993. Primary structure and cell-type specific expression of a gustatory G protein-coupled receptor related to olfactory receptors. J. Bio. Chem.). This provides an addition example of receptors

which can be isolated, expressed in minicells and then be used for identification of specific substances in various matrices in similar manners as identified for olfactory receptor minicells.

[0943] As a non-limiting example of minicell use in toxicological/environmental detection, arrays could be constructed in which each well contains a distinct minicell subtype displaying membrane-bound proteins or other molecules for each of several potential toxins or agents in the environment. For example, minicells in such a format could be used to determine which agents are present in the environment as a consequence of a chemical and/or biological weapons attack. Non-limiting examples of biosensors that have been used toxicological/environmental detection include those described by Sticher et al., Development and characterization of a whole-cell bioluminescent sensor for bioavailable middle-chain alkanes in contaminated ground-water samples, Appl. Envir. Microbiol. 63:4053-4060, 1997; Willardson et al., Development and Testing of a Bacterial Biosensor for Toluene-Based Environmental Contaminants, Appl. Envir. Microbiol. 64:1006-1012, 1998; Lars et al., Detection of Oxytetracycline Production by *Streptomyces rimosus* in Soil Microcosms by Combining Whole-Cell Biosensors and Flow Cytometry, Appl. Envir. Microbiol. 67:239-244, 2001; Højberg et al., Oxygen-Sensing Reporter Strain of *Pseudomonas fluorescens* for Monitoring the Distribution of Low-Oxygen Habitats in Soil, Appl. Envir. Microbiol. 1999 65: 4085-4093, 1999; R. P. Hollis et al., Design and Application of a Biosensor for Monitoring Toxicity of Compounds to Eukaryotes, Appl. Envir. Microbiol. 66: 1676-1679, 2000; Heitzer et al., Optical biosensor for environmental on-line monitoring of naphthalene and salicylate bioavailability with an immobilized bioluminescent catabolic reporter bacterium, Appl. Envir. Microbiol. 60:1487-1494, 1994; Selifonova et al., Bioluminescent sensors for detection of bioavailable Hg(II) in the environment, Appl. Envir. Microbiol. 59: 3083-3090, 1993; Jaeger et al., Mapping of Sugar and Amino Acid Availability in Soil around Roots with Bacterial Sensors of Sucrose and Tryptophan, Appl. Envir. Microbiol. 65: 2685-2690, 1999; and Larsen et al., A Microsensor for Nitrate Based on Immobilized Denitrifying Bacteria, Appl. Envir. Microbiol. 62: 1248-1251, 1996.

[0944] XVIII.C. Toxin Elimination

[0945] In another embodiment of the invention, minicells displaying a receptor for a particular toxic agent could be used for the elimination of the agent from the environment. In a non-limiting example of this technology, minicells could be placed in a filtering apparatus to eliminate the toxic agent from the environment (e.g., air, water soil). In the example of atmospheric contamination, the air would be circulated through a forced air system containing in-line filters composed of a housing, support matrix and receptor/minicells. As air passes over the minicells, the toxin is bound to the receptor. The purified air passed out of the system and into the atmosphere. A similar method for water purification would follow a similar protocol replacing the receptor for the toxin with the receptor or other protein binding a unique epitope on contaminant wishing to be removed. Examples include but are not limited to removing toxins, parasites or microbes from the matrix such as water or air. This represent non-limiting example of minicell-based technology for expression of functional receptors or binding moieties of

receptors on the minicell's surface for the specific purpose of selectively capturing, identifying, quantifying and/or removing molecules of interest for environmental compartments to include but not limited to air water, soil, other gas phases or liquid solutions.

[0946] Representative toxins include, but are not limited to, those associated with "red tides"; eubacterial toxins, such as those toxins produced by *Corynebacterium diphtheriae* (diphtheria), *Bordetella pertussis* (whooping cough), *Vibrio cholerae* (cholera), *Bacillus anthracis* (anthrax), *Clostridium botulinum* (botulism), *Clostridium tetani* (tetanus), and enterohemorrhagic *Escherichia coli* (bloody diarrhea and hemolytic uremic syndrome); and fungal toxins (e.g., aflatoxin, gliotoxin, cyclopeptides, orellanine, gyrometrin, coprine, muscarine, ibotenic acid, psilocybin, psilocin and baeocystin).

[0947] The treatment of "red tides" with minicells exemplifies this aspect of the invention. A red tide occurs as a result of a higher-than-normal concentration of an algae or dinoflagellate which, when present in dense concentrations as a result of a "bloom," form colored patches on the surface of water. The colored patches are pink, violet, orange, yellow, blue, green, brown, or red, with red being the most common color. The organisms that cause red tides often produce toxins that have negative impacts on other organisms, including humans.

[0948] For example, *Karenia brevis* (formerly *Gymnodinium breve*) produces a toxin (domoic acid) that affects the central nervous system of fish, shellfish and other organisms, resulting in a state of paralysis. Alexandrium species (e.g., *A. tamarense*, *A. fundyense*), Dinophysis and Gonyaulax species; and Pseudo-nitzschia multiseriata, which cause, respectively, paralytic, diarrhetic and amnesic shellfish poisoning. Because shellfish containing the toxin taste and appear the same as shellfish that do not, and cooking does not destroy the toxin, human ingestion of the former can cause disease in humans and other organisms. For example, one form of paralytic shellfish poisoning, which can be fatal to humans, results from saxitoxin, which is produced by *Gonyaulax tamarensis*, *Protogonyaulax catenella*, and other species. Other algae that can result in red tides include *Gonyaulax catenella*, and *Ptychodiscus brevis*.

[0949] Minicells that comprise a binding moiety of an organism that produces a red tide, or of the toxin produced thereby, can be used for remediation. For example, a minicell having a binding moiety directed to a red tide-producing organism can be used to deliver an antibiotic thereto, and a minicell with a binding moiety directed to a toxin can be used to bind and/or internalize the toxin. As is explained in more detail elsewhere herein, a minicell with a binding moiety directed to a toxin can also be used for therapeutic purposes.

#### [0950] XVIII.D. Bioremediation

[0951] In another non-limiting example of the potential use of minicells in a toxicological context is their use in bioremediation, the process by which living organisms act to degrade or transform hazardous organic contaminants. As used herein, "bioremediation" is the process of using biological or biologically derived compositions that alter the chemical structure and/or bind, an undesirable substance in order to reduce the effective concentration of the undesirable

substance, thereby reducing or eliminating the effect(s) of the undesirable compound on the environment. Undesirable substances include, but are not limited to, pollutants (e.g., heavy metals, pesticides, herbicides, petroleum products); biological toxins (e.g., such as those produced by "red tides", e.g., domoic acid, saxitoxin); pathogens (e.g., viruses, eubacteria); organisms that produce toxins; biological waste products (e.g., sewage, guano), and undesirable organisms therewithin (e.g., pathogenic eubacteria).

[0952] The term "bioremediation" encompasses both biodegradation, the breakdown of organic substances by microorganisms, and biotransformation, the alteration of the structure of a compound by a living organism or enzyme. The minicells of the invention may be incorporated into biofilters, i.e., devices in which gases, liquids, powders and the like are passed through media containing biodegrading minicells, including but not limited to devices that biodegrade volatile organic compounds in air by passing the air therethrough.

[0953] Bioremediation can be used to process undesirable substances in a composition prior to or after the release of the composition into the environment. For example, bioremediation can be applied in sewage treatment plants to process sewage prior to its release, or to sewage that has been accidentally or otherwise released into the environment.

[0954] Environmental microbiologists have sought to identify and use specific bacteria that degrade pollutants and other environmental contaminants. See, for example, Chakrabarty, Microbial Degradation of Toxic Chemicals: Evolutionary Insights and Practical Considerations, Am. Soc. Micro. Biol. News 62:130-137, 1996; and U.S. Pat. Nos. 4,511,657; 4,493,895; 4,871,673; and 4,535,061. In instances where a live organism is placed into the environment to process undesirable substances, there is a concern that the organism might have undesirable effects that would be made more deleterious due to the ability of the live organism to replicate (Sayler G S, Ripp S. Field applications of genetically engineered microorganisms for bioremediation processes. Curr Opin Biotechnol. 2000 June;11(3):286-9; and Diaz E, Ferrandez A, Prieto M A, Garcia J L. Biodegradation of aromatic compounds by *Escherichia coli*. Microbiol Mol Biol Rev. 2001 December;65(4):523-69). For example, when it has been proposed to use genetically altered eubacteria to process oil spills, the concern has been raised that the eubacteria might spread beyond the oil spill and into supplies of petroleum products that are used to produce energy, where they would process and render useless the stored petroleum products. However, because they lack the ability to replicate, such a scenario will not occur when minicells are used for bioremediation.

[0955] By way of non-limiting example, octane enhances such as methyl t-butyl ether or aromatic hydrocarbons contaminate the aquifer and soil. These agents negatively impact the many microbes in the effected area thus limiting capability of the microbial community rectify the environmental insult. Bioaugmentation, the addition to the environment of microorganisms that can metabolize and grow on specific organic compounds, to facilitate degradation may prove useful, but concerns exist relative to the regulation of newly introduced bacteria. The minicell provides a vehicle to accomplish biodegradation without bacterial overgrowth.

[0956] Diphenyl ethers and cyclic ethers such as dioxane and furan have shown to be metabolized by soil bacteria. Using classic isolation and screening techniques identified above, genes encoding for the oxygenases or hydrolyases are isolated. The enzyme sequence is inserted into an expression vector using standard molecular biology techniques. Minicell producing bacteria are transformed with these vectors and cultured. Minicells are isolated from the culture as previously described and subsequently induced. The minicells are applied to the area contaminated with aromatic hydrocarbons. These compounds are transported either actively or passively in to the minicell and subsequently degraded by the oxygenase or hydrolyase. One advantage of this focused degradation is the minimizing of feedback inhibition because the only machinery of consequence in the minicell is that related to the degradation of the ether compounds.

[0957] Similarly, beginning with genetic material from *Dehalobacter* enzymes responsible for the biodegrading of tetrachloroethane could be isolated as described above. The sequence for the enzyme is inserted into the expression vector and used to transform minicell-producing bacteria. The bacteria are cultured, minicells isolated from the culture and the minicells induced as previously described. Minicell preps are lyophilized using standard lyophilization techniques. The resulting material is transported to the site of tetrachloroethene contamination and reconstituted and applied. As the tetrachloroethene was assimilated, it is degraded by the enzyme system.

[0958] These are non-limiting examples scope of bioremediation/biotransformation using minicell technology. The scope of the invention includes taking advantage of metabolic pathways organism in general to include but not limited to eukaryotes, prokaryotes, fungi, animals or plants.

#### [0959] XVIII.E. Fermentation

[0960] Delivery of specific enzymes in an untargeted fashion by the minicell allows for packaged delivery without the increased biomass and complex metabolic products associated with processes using live organisms. This aspect can be taken advantage of in fermentation, where the addition of minicells into which unique enzymes have been added are used to modulate the composition of the environment to include but not limited to the alcohol, sugar and acid levels.

#### [0961] XVIII.F. Pesticides

[0962] *Bacillus thuringiensis* produces a toxin that kills plant chewing insect larvae as well as mosquito larvae. The toxin, Cry1Ac, binds to aminopeptidase N receptor on the endothelium of the midgut. Minicell technology is allows for delivery of the toxin. The toxin sequence is modified by ligation of a sequence coding for a transmembrane domain as previously described. The sequence for this fusion protein inserted into an expression vector using standard molecular biology techniques. To facilitate the consumption of the toxin/minicell plasmids containing sequences incorporating the sequence for pheromones coupled at the C-terminus to the sequence for a transmembrane domain is generated using standard molecular biological techniques. This fusion protein sequence is inserted into the expression containing coding region for the toxin fusion protein or inserted into a unique expression vector. Minicell producing bacteria are

transformed with these vectors and cultured. Minicells are isolated from the culture as previously described and subsequently induced. The minicells are distributed (e.g crop dusting) to the area of infestation. The toxin/minicells are ingested by the larvae and kill the larvae as the minicells passes through the gut.

#### [0963] XIX. Pharmaceutical Compositions

[0964] Another aspect of the invention is drawn to compositions, including but not limited to pharmaceutical compositions. According to the invention, a "composition" refers to a mixture comprising at least one carrier, preferably a physiologically acceptable carrier, and one or more minicell compositions. The term "carrier" defines a chemical compound that does not inhibit or prevent the incorporation of the biologically active peptide(s) into cells or tissues. A carrier typically is an inert substance that allows an active ingredient to be formulated or compounded into a suitable dosage form (e.g., a pill, a capsule, a gel, a film, a tablet, a microparticle (e.g., a microsphere), a solution; an ointment; a paste, an aerosol, a droplet, a colloid or an emulsion etc.). A "physiologically acceptable carrier" is a carrier suitable for use under physiological conditions that does not abrogate (reduce, inhibit, or prevent) the biological activity and properties of the compound. For example, dimethyl sulfoxide (DMSO) is a carrier as it facilitates the uptake of many organic compounds into the cells or tissues of an organism. Preferably, the carrier is a physiologically acceptable carrier, preferably a pharmaceutically or veterinarily acceptable carrier, in which the minicell composition is disposed.

[0965] A "pharmaceutical composition" refers to a composition wherein the carrier is a pharmaceutically acceptable carrier, while a "veterinary composition" is one wherein the carrier is a veterinarily acceptable carrier. The term "pharmaceutically acceptable carrier" or "veterinarily acceptable carrier" includes any medium or material that is not biologically or otherwise undesirable, i.e., the carrier may be administered to an organism along with a minicell composition without causing any undesirable biological effects or interacting in a deleterious manner with the complex or any of its components or the organism. Examples of pharmaceutically acceptable reagents are provided in The United States Pharmacopeia, The National Formulary, United States Pharmacopeial Convention, Inc., Rockville, Md. 1990, hereby incorporated by reference herein into the present application. The terms "therapeutically effective amount" or "pharmaceutically effective amount" mean an amount sufficient to induce or effectuate a measurable response in the target cell, tissue, or body of an organism. What constitutes a therapeutically effective amount will depend on a variety of factors, which the knowledgeable practitioner will take into account in arriving at the desired dosage regimen.

[0966] The compositions of the invention can further comprise other chemical components, such as diluents and excipients. A "diluent" is a chemical compound diluted in a solvent, preferably an aqueous solvent, that facilitates dissolution of the composition in the solvent, and it may also serve to stabilize the biologically active form of the composition or one or more of its components. Salts dissolved in buffered solutions are utilized as diluents in the art. For example, preferred diluents are buffered solutions containing one or more different salts. A preferred buffered solution is phosphate buffered saline (particularly in conjunction with

compositions intended for pharmaceutical administration), as it mimics the salt conditions of human blood. Since buffer salts can control the pH of a solution at low concentrations, a buffered diluent rarely modifies the biological activity of a biologically active peptide.

[0967] An “excipient” is any more or less inert substance that can be added to a composition in order to confer a suitable property, for example, a suitable consistency or to form a drug. Suitable excipients and carriers include, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol cellulose preparations such as, for example, maize starch, wheat starch, rice starch, agar, pectin, xanthan gum, guar gum, locust bean gum, hyaluronic acid, casein potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, polyacrylate, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents can also be included, such as cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate. Other suitable excipients and carriers include hydrogels, gellable hydrocolloids, and chitosan. Chitosan microspheres and microcapsules can be used as carriers. See WO 98/52547 (which describes microsphere formulations for targeting compounds to the stomach, the formulations comprising an inner core (optionally including a gelled hydrocolloid) containing one or more active ingredients, a membrane comprised of a water insoluble polymer (e.g., ethylcellulose) to control the release rate of the active ingredient(s), and an outer layer comprised of a bioadhesive cationic polymer, for example, a cationic polysaccharide, a cationic protein, and/or a synthetic cationic polymer; U.S. Pat. No. 4,895,724. Typically, chitosan is cross-linked using a suitable agent, for example, glutaraldehyde, glyoxal, epichlorohydrin, and succinaldehyde. Compositions employing chitosan as a carrier can be formulated into a variety of dosage forms, including pills, tablets, microparticles, and microspheres, including those providing for controlled release of the active ingredient(s). Other suitable bioadhesive cationic polymers include acidic gelatin, polygalactosamine, polyamino acids such as polylysine, polyhistidine, polyornithine” polyquaternary compounds, prolamine, polyimine, diethylaminoethyl-dextran (DEAE), DEAE-imine, DEAE-methacrylate, DEAE-acrylamide, DEAE-dextran, DEAE-cellulose, poly-p-aminostyrene, polyoxethane, copolymethacrylates, polyamidoamines, cationic starches, polyvinylpyridine, and polythiodiethylaminomethylethylene.

[0968] The compositions of the invention can be formulated in any suitable manner. Minicell compositions may be uniformly (homogeneously) or non-uniformly (heterogeneously) dispersed in the carrier. Suitable formulations include dry and liquid formulations. Dry formulations include freeze dried and lyophilized powders, which are particularly well suited for aerosol delivery to the sinuses or lung, or for long term storage followed by reconstitution in a suitable diluent prior to administration. Other preferred dry formulations include those wherein a composition according to the invention is compressed into tablet or pill form suitable for oral administration or compounded into a sustained release formulation. When the composition is intended for oral administration but is to be delivered to epithelium in the intestines, it is preferred that the formulation be encapsulated with an enteric coating to protect the formulation and prevent premature release of the minicell

compositions included therein. As those in the art will appreciate, the compositions of the invention can be placed into any suitable dosage form. Pills and tablets represent some of such dosage forms. The compositions can also be encapsulated into any suitable capsule or other coating material, for example, by compression, dipping, pan coating, spray drying, etc. Suitable capsules include those made from gelatin and starch. In turn, such capsules can be coated with one or more additional materials, for example, and enteric coating, if desired. Liquid formulations include aqueous formulations, gels, and emulsions.

[0969] Some preferred embodiments concern compositions that comprise a bioadhesive, preferably a mucoadhesive, coating. A “bioadhesive coating” is a coating that allows a substance (e.g., a minicell composition) to adhere to a biological surface or substance better than occurs absent the coating. A “mucoadhesive coating” is a preferred bioadhesive coating that allows a substance, for example, a composition according to the invention, to adhere better to mucosa occurs absent the coating. For example, micronized particles (e.g., particles having a mean diameter of about 5, 10, 25, 50, or 100  $\mu\text{m}$ ) can be coated with a mucoadhesive. The coated particles can then be assembled into a dosage form suitable for delivery to an organism. Preferably, and depending upon the location where the cell surface transport moiety to be targeted is expressed, the dosage form is then coated with another coating to protect the formulation until it reaches the desired location, where the mucoadhesive enables the formulation to be retained while the composition interacts with the target cell surface transport moiety.

[0970] The compositions of the invention may be administered to any organism, preferably an animal, preferably a mammal, bird, fish, insect, or arachnid. Preferred mammals include bovine, canine, equine, feline, ovine, and porcine animals, and non-human primates. Humans are particularly preferred. Multiple techniques of administering or delivering a compound exist in the art including, but not limited to, oral, rectal (e.g. an enema or suppository) aerosol (e.g., for nasal or pulmonary delivery), parenteral, and topical administration. Preferably, sufficient quantities of the biologically active peptide are delivered to achieve the intended effect. The particular amount of composition to be delivered will depend on many factors, including the effect to be achieved, the type of organism to which the composition is delivered, delivery route, dosage regimen, and the age, health, and sex of the organism. As such, the particular dosage of a composition incorporated into a given formulation is left to the ordinarily skilled artisan’s discretion.

[0971] Those skilled in the art will appreciate that when the compositions of the present invention are administered as agents to achieve a particular desired biological result, which may include a therapeutic or protective effect(s) (including vaccination), it may be necessary to combine the fusion proteins of the invention with a suitable pharmaceutical carrier. The choice of pharmaceutical carrier and the preparation of the fusion protein as a therapeutic or protective agent will depend on the intended use and mode of administration. Suitable formulations and methods of administration of therapeutic agents include those for oral, pulmonary, nasal, buccal, ocular, dermal, rectal, or vaginal delivery.

[0972] Depending on the mode of delivery employed, the context-dependent functional entity can be delivered in a

variety of pharmaceutically acceptable forms. For example, the context-dependent functional entity can be delivered in the form of a solid, solution, emulsion, dispersion, micelle, liposome, and the like, incorporated into a pill, capsule, tablet, suppository, aerosol, droplet, or spray. Pills, tablets, suppositories, aerosols, powders, droplets, and sprays may have complex, multilayer structures and have a large range of sizes. Aerosols, powders, droplets, and sprays may range from small (1 micron) to large (200 micron) in size.

[0973] Pharmaceutical compositions of the present invention can be used in the form of a solid, a lyophilized powder, a solution, an emulsion, a dispersion, a micelle, a liposome, and the like, wherein the resulting composition contains one or more of the compounds of the present invention, as an active ingredient, in admixture with an organic or inorganic carrier or excipient suitable for enteral or parenteral applications. The active ingredient may be compounded, for example, with the usual non-toxic, pharmaceutically acceptable carriers for tablets, pellets, capsules, suppositories, solutions, emulsions, suspensions, and any other form suitable for use. The carriers which can be used include glucose, lactose, mannose, gum acacia, gelatin, mannitol, starch paste, magnesium trisilicate, talc, corn starch, keratin, colloidal silica, potato starch, urea, medium chain length triglycerides, dextrans, and other carriers suitable for use in manufacturing preparations, in solid, semisolid, or liquid form. In addition auxiliary, stabilizing, thickening and coloring agents and perfumes may be used. Examples of a stabilizing dry agent includes triulose, preferably at concentrations of 0.1% or greater (See, e.g., U.S. Pat. No. 5,314,695). The active compound is included in the pharmaceutical composition in an amount sufficient to produce the desired effect upon the process or condition of diseases.

#### [0974] XX. Small Molecules

[0975] The term "small molecule" includes any chemical or other moiety that can act to affect biological processes. Small molecules can include any number of therapeutic agents presently known and used, or can be small molecules synthesized in a library of such molecules for the purpose of screening for biological function(s). Small molecules are distinguished from macromolecules by size. The small molecules of this invention usually have molecular weight less than about 5,000 daltons (Da), preferably less than about 2,500 Da, more preferably less than 1,000 Da, most preferably less than about 500 Da.

[0976] Small molecules include without limitation organic compounds, peptidomimetics and conjugates thereof. As used herein, the term "organic compound" refers to any carbon-based compound other than macromolecules such nucleic acids and polypeptides. In addition to carbon, organic compounds may contain calcium, chlorine, fluorine, copper, hydrogen, iron, potassium, nitrogen, oxygen, sulfur and other elements. An organic compound may be in an aromatic or aliphatic form. Non-limiting examples of organic compounds include acetones, alcohols, anilines, carbohydrates, monosaccharides, oligosaccharides, polysaccharides, amino acids, nucleosides, nucleotides, lipids, retinoids, steroids, proteoglycans, ketones, aldehydes, saturated, unsaturated and polyunsaturated fats, oils and waxes, alkenes, esters, ethers, thiols, sulfides, cyclic compounds, heterocyclic compounds, imidizoles and phenols. An organic compound as used herein also includes nitrated

organic compounds and halogenated (e.g., chlorinated) organic compounds. Methods for preparing peptidomimetics are described below. Collections of small molecules, and small molecules identified according to the invention are characterized by techniques such as accelerator mass spectrometry (AMS; see Turteltaub et al., *Curr Pharm Des* 2000 6(10):991-1007, Bioanalytical applications of accelerator mass spectrometry for pharmaceutical research; and Enjalbal et al., *Mass Spectrom Rev* 2000 19(3):139-61, Mass spectrometry in combinatorial chemistry.)

[0977] Preferred small molecules are relatively easier and less expensively manufactured, formulated or otherwise prepared. Preferred small molecules are stable under a variety of storage conditions. Preferred small molecules may be placed in tight association with macromolecules to form molecules that are biologically active and that have improved pharmaceutical properties. Improved pharmaceutical properties include changes in circulation time, distribution, metabolism, modification, excretion, secretion, elimination, and stability that are favorable to the desired biological activity. Improved pharmaceutical properties include changes in the toxicological and efficacy characteristics of the chemical entity.

#### [0978] XXI. Polypeptides and Derivatives

##### [0979] XXI.A. Polypeptides

[0980] As used herein, the term "polypeptide" includes proteins, fusion proteins, oligopeptides and polypeptide derivatives, with the exception that peptidomimetics are considered to be small molecules herein. Although they are polypeptides, antibodies and their derivatives are described in a separate section. Antibodies and antibody derivatives are described in a separate section, but antibodies and antibody derivatives are, for purposes of the invention, treated as a subclass of the polypeptides and derivatives.

[0981] A "protein" is a molecule having a sequence of amino acids that are linked to each other in a linear molecule by peptide bonds. The term protein refers to a polypeptide that is isolated from a natural source, or produced from an isolated cDNA using recombinant DNA technology; and has a sequence of amino acids having a length of at least about 200 amino acids.

[0982] A "fusion protein" is a type of recombinant protein that has an amino acid sequence that results from the linkage of the amino acid sequences of two or more normally separate polypeptides.

[0983] A "protein fragment" is a proteolytic fragment of a larger polypeptide, which may be a protein or a fusion protein. A proteolytic fragment may be prepared by *in vivo* or *in vitro* proteolytic cleavage of a larger polypeptide, and is generally too large to be prepared by chemical synthesis. Proteolytic fragments have amino acid sequences having a length from about 200 to about 1,000 amino acids.

[0984] An "oligopeptide" is a polypeptide having a short amino acid sequence (i.e., 2 to about 200 amino acids). An oligopeptide is generally prepared by chemical synthesis.

[0985] Although oligopeptides and protein fragments may be otherwise prepared, it is possible to use recombinant DNA technology and/or *in vitro* biochemical manipulations. For example, a nucleic acid encoding an amino acid sequence may be prepared and used as a template for *in vitro*

transcription/translation reactions. In such reactions, an exogenous nucleic acid encoding a preselected polypeptide is introduced into a mixture that is essentially depleted of exogenous nucleic acids that contains all of the cellular components required for transcription and translation. One or more radiolabeled amino acids are added before or with the exogenous DNA, and transcription and translation are allowed to proceed. Because the only nucleic acid present in the reaction mix is the exogenous nucleic acid added to the reaction, only polypeptides encoded thereby are produced, and incorporate the radiolabelled amino acid(s). In this manner, polypeptides encoded by a preselected exogenous nucleic acid are radiolabeled. Although other proteins are present in the reaction mix, the preselected polypeptide is the only one that is produced in the presence of the radiolabeled amino acids and is thus uniquely labeled.

[0986] As is explained in detail below, "polypeptide derivatives" include without limitation mutant polypeptides, chemically modified polypeptides, and peptidomimetics.

[0987] The polypeptides of this invention, including the analogs and other modified variants, may generally be prepared following known techniques. Preferably, synthetic production of the polypeptide of the invention may be according to the solid phase synthetic method. For example, the solid phase synthesis is well understood and is a common method for preparation of polypeptides, as are a variety of modifications of that technique [Merrifield (1964), *J. Am. Chem. Soc.*, 85: 2149; Stewart and Young (1984), *Solid Phase polypeptide Synthesis*, Pierce Chemical Company, Rockford, Ill.; Bodansky and Bodansky (1984), *The Practice of polypeptide Synthesis*, Springer-Verlag, New York; Atherton and Sheppard (1989), *Solid Phase polypeptide Synthesis: A Practical Approach*, IRL Press, New York]. See, also, the specific method described in Example 1 below.

[0988] Alternatively, polypeptides of this invention may be prepared in recombinant systems using polynucleotide sequences encoding the polypeptides. For example, fusion proteins are typically prepared using recombinant DNA technology.

[0989] XXI.B. Polypeptide Derivatives

[0990] A "derivative" of a polypeptide is a compound that is not, by definition, a polypeptide, i.e., it contains at least one chemical linkage that is not a peptide bond. Thus, polypeptide derivatives include without limitation proteins that naturally undergo post-translational modifications such as, e.g., glycosylation. It is understood that a polypeptide of the invention may contain more than one of the following modifications within the same polypeptide. Preferred polypeptide derivatives retain a desirable attribute, which may be biological activity; more preferably, a polypeptide derivative is enhanced with regard to one or more desirable attributes, or has one or more desirable attributes not found in the parent polypeptide. Although they are described in this section, peptidomimetics are taken as small molecules in the present disclosure.

[0991] XXI.C. Mutant Polypeptide Derivatives

[0992] A polypeptide having an amino acid sequence identical to that found in a protein prepared from a natural source is a "wildtype" polypeptide. Mutant oligopeptides can be prepared by chemical synthesis, including without limitation combinatorial synthesis. Mutant polypeptides

larger than oligopeptides can be prepared using recombinant DNA technology by altering the nucleotide sequence of a nucleic acid encoding a polypeptide. Although some alterations in the nucleotide sequence will not alter the amino acid sequence of the polypeptide encoded thereby ("silent" mutations), many will result in a polypeptide having an altered amino acid sequence that is altered relative to the parent sequence. Such altered amino acid sequences may comprise substitutions, deletions and additions of amino acids, with the proviso that such amino acids are naturally occurring amino acids.

[0993] Thus, subjecting a nucleic acid that encodes a polypeptide to mutagenesis is one technique that can be used to prepare mutant polypeptides, particularly ones having substitutions of amino acids but no deletions or insertions thereof. A variety of mutagenic techniques are known that can be used in vitro or in vivo including without limitation chemical mutagenesis and PCR-mediated mutagenesis. Such mutagenesis may be randomly targeted (i.e., mutations may occur anywhere within the nucleic acid) or directed to a section of the nucleic acid that encodes a stretch of amino acids of particular interest. Using such techniques, it is possible to prepare randomized, combinatorial or focused compound libraries, pools and mixtures.

[0994] Polypeptides having deletions or insertions of naturally occurring amino acids may be synthetic oligopeptides that result from the chemical synthesis of amino acid sequences that are based on the amino acid sequence of a parent polypeptide but which have one or more amino acids inserted or deleted relative to the sequence of the parent polypeptide. Insertions and deletions of amino acid residues in polypeptides having longer amino acid sequences may be prepared by directed mutagenesis.

[0995] XXI.D. Chemically Modified Polypeptides

[0996] As contemplated by this invention, the term "polypeptide" includes those having one or more chemical modification relative to another polypeptide, i.e., chemically modified polypeptides. The polypeptide from which a chemically modified polypeptide is derived may be a wild-type protein, a mutant protein or a mutant polypeptide, or polypeptide fragments thereof; an antibody or other polypeptide ligand according to the invention including without limitation single-chain antibodies, bacterial proteins and polypeptide derivatives thereof; or polypeptide ligands prepared according to the disclosure. Preferably, the chemical modification(s) confer(s) or improve(s) desirable attributes of the polypeptide but does not substantially alter or compromise the biological activity thereof. Desirable attributes include but are limited to increased shelf-life; enhanced serum or other in vivo stability; resistance to proteases; and the like. Such modifications include by way of non-limiting example N-terminal acetylation, glycosylation, and biotinylation.

[0997] XXI.D.1. Polypeptides with N-Terminal or C-Terminal Chemical Groups

[0998] An effective approach to confer resistance to peptidases acting on the N-terminal or C-terminal residues of a polypeptide is to add chemical groups at the polypeptide termini, such that the modified polypeptide is no longer a substrate for the peptidase. One such chemical modification is glycosylation of the polypeptides at either or both termini.

Certain chemical modifications, in particular N-terminal glycosylation, have been shown to increase the stability of polypeptides in human serum (Powell et al. (1993), *Pharma. Res.* 10: 1268-1273). Other chemical modifications which enhance serum stability include, but are not limited to, the addition of an N-terminal alkyl group, consisting of a lower alkyl of from 1 to 20 carbons, such as an acetyl group, and/or the addition of a C-terminal amide or substituted amide group.

[0999] XXI.D.2. Polypeptides with a Terminal D-Amino Acid

[1000] The presence of an N-terminal D-amino acid increases the serum stability of a polypeptide that otherwise contains L-amino acids, because exopeptidases acting on the N-terminal residue cannot utilize a D-amino acid as a substrate. Similarly, the presence of a C-terminal D-amino acid also stabilizes a polypeptide, because serum exopeptidases acting on the C-terminal residue cannot utilize a D-amino acid as a substrate. With the exception of these terminal modifications, the amino acid sequences of polypeptides with N-terminal and/or C-terminal D-amino acids are usually identical to the sequences of the parent L-amino acid polypeptide.

[1001] XXI.D.3. Polypeptides with Substitution of Natural Amino Acids by Unnatural Amino Acids

[1002] Substitution of unnatural amino acids for natural amino acids in a subsequence of a polypeptide can confer or enhance desirable attributes including biological activity. Such a substitution can, for example, confer resistance to proteolysis by exopeptidases acting on the N-terminus. The synthesis of polypeptides with unnatural amino acids is routine and known in the art (see, for example, Coller, et al. (1993), cited above).

[1003] XXI.D.4. Post-Translational Chemical Modifications

[1004] Different host cells will contain different post-translational modification mechanisms that may provide particular types of post-translational modification of a fusion protein if the amino acid sequences required for such modifications is present in the fusion protein. A large number (~100) of post-translational modifications have been described, a few of which are discussed herein. One skilled in the art will be able to choose appropriate host cells, and design chimeric genes that encode protein members comprising the amino acid sequence needed for a particular type of modification.

[1005] Glycosylation is one type of post-translational chemical modification that occurs in many eukaryotic systems, and may influence the activity, stability, pharmacogenetics, immunogenicity and/or antigenicity of proteins. However, specific amino acids must be present at such sites to recruit the appropriate glycosylation machinery, and not all host cells have the appropriate molecular machinery. *Saccharomyces cerevisiae* and *Pichia pastoris* provide for the production of glycosylated proteins, as do expression systems that utilize insect cells, although the pattern of glycosylation may vary depending on which host cells are used to produce the fusion protein.

[1006] Another type of post-translation modification is the phosphorylation of a free hydroxyl group of the side chain

of one or more Ser, Thr or Tyr residues. Protein kinases catalyze such reactions. Phosphorylation is often reversible due to the action of a protein phosphatase, an enzyme that catalyzes the dephosphorylation of amino acid residues.

[1007] Differences in the chemical structure of amino terminal residues result from different host cells, each of which may have a different chemical version of the methionine residue encoded by a start codon, and these will result in amino termini with different chemical modifications.

[1008] For example, many or most bacterial proteins are synthesized with an amino terminal amino acid that is a modified form of methionine, i.e., N-formyl-methionine (fMet). Although the statement is often made that all bacterial proteins are synthesized with an fMet initiator amino acid; although this may be true for *E. coli*, recent studies have shown that it is not true in the case of other bacteria such as *Pseudomonas aeruginosa* (Newton et al., *J. Biol. Chem.* 274:22143-22146, 1999). In any event, in *E. coli*, the formyl group of fMet is usually enzymatically removed after translation to yield an amino terminal methionine residue, although the entire fMet residue is sometimes removed (see Hershey, Chapter 40, "Protein Synthesis" in: *Escherichia Coli and Salmonella Typhimurium: Cellular and Molecular Biology*, Neidhardt, Frederick C., Editor in Chief, American Society for Microbiology, Washington, D.C., 1987, Volume 1, pages 613-647, and references cited therein.) *E. coli* mutants that lack the enzymes (such as, e.g., formylase) that catalyze such post-translational modifications will produce proteins having an amino terminal fMet residue (Guillon et al., *J. Bacteriol.* 174:4294-4301, 1992).

[1009] In eukaryotes, acetylation of the initiator methionine residue, or the penultimate residue if the initiator methionine has been removed, typically occurs co- or post-translationally. The acetylation reactions are catalyzed by N-terminal acetyltransferases (NATs, a.k.a. N-alpha-acetyltransferases), whereas removal of the initiator methionine residue is catalyzed by methionine aminopeptidases (for reviews, see Bradshaw et al., *Trends Biochem. Sci.* 23:263-267, 1998; and Driessen et al., *CRC Crit. Rev. Biochem.* 18:281-325, 1985). Amino terminally acetylated proteins are said to be "N-acetylated," "N alpha acetylated" or simply "acetylated."

[1010] Another post-translational process that occurs in eukaryotes is the alpha-amidation of the carboxy terminus. For reviews, see Eipper et al. *Annu. Rev. Physiol.* 50:333-344, 1988, and Bradbury et al. *Lung Cancer* 14:239-251, 1996. About 50% of known endocrine and neuroendocrine peptide hormones are alpha-amidated (Treston et al., *Cell Growth Differ.* 4:911-920, 1993). In most cases, carboxy alpha-amidation is required to activate these peptide hormones.

[1011] XXI.E. Peptidomimetics

[1012] In general, a polypeptide mimetic ("peptidomimetic") is a molecule that mimics the biological activity of a polypeptide but is no longer peptidic in chemical nature. By strict definition, a peptidomimetic is a molecule that contains no peptide bonds (that is, amide bonds between amino acids). However, the term peptidomimetic is sometimes used to describe molecules that are no longer completely peptidic in nature, such as pseudo-peptides, semi-peptides and peptoids. Examples of some peptidomimetics



by the broader definition (where part of a polypeptide is replaced by a structure lacking peptide bonds) are described below. Whether completely or partially non-peptide, peptidomimetics according to this invention provide a spatial arrangement of reactive chemical moieties that closely resembles the three-dimensional arrangement of active groups in the polypeptide on which the peptidomimetic is based. As a result of this similar active-site geometry, the peptidomimetic has effects on biological systems that are similar to the biological activity of the polypeptide.

[1013] There are several potential advantages for using a mimetic of a given polypeptide rather than the polypeptide itself. For example, polypeptides may exhibit two undesirable attributes, i.e., poor bioavailability and short duration of action. Peptidomimetics are often small enough to be both orally active and to have a long duration of action. There are also problems associated with stability, storage and immunoreactivity for polypeptides that are not experienced with peptidomimetics.

[1014] Candidate, lead and other polypeptides having a desired biological activity can be used in the development of peptidomimetics with similar biological activities. Techniques of developing peptidomimetics from polypeptides are known. Peptide bonds can be replaced by non-peptide bonds that allow the peptidomimetic to adopt a similar structure, and therefore biological activity, to the original polypeptide. Further modifications can also be made by replacing chemical groups of the amino acids with other chemical groups of similar structure. The development of peptidomimetics can be aided by determining the tertiary structure of the original polypeptide, either free or bound to a ligand, by NMR spectroscopy, crystallography and/or computer-aided molecular modeling. These techniques aid in the development of novel compositions of higher potency and/or greater bioavailability and/or greater stability than the original polypeptide (Dean (1994), *BioEssays*, 16: 683-687; Cohen and Shatzmiller (1993), *J. Mol. Graph.*, 11: 166-173; Wiley and Rich (1993), *Med. Res. Rev.*, 13: 327-384; Moore (1994), *Trends Pharmacol. Sci.*, 15: 124-129; Hruby (1993), *Biopolymers*, 33: 1073-1082; Bugg et al. (1993), *Sci. Am.*, 269: 92-98, all incorporated herein by reference).

[1015] Thus, through use of the methods described above, the present invention provides compounds exhibiting enhanced therapeutic activity in comparison to the polypeptides described above. The peptidomimetic compounds obtained by the above methods, having the biological activity of the above named polypeptides and similar three-dimensional structure, are encompassed by this invention. It will be readily apparent to one skilled in the art that a peptidomimetic can be generated from any of the modified polypeptides described in the previous section or from a polypeptide bearing more than one of the modifications described from the previous section. It will furthermore be apparent that the peptidomimetics of this invention can be further used for the development of even more potent non-peptidic compounds, in addition to their utility as therapeutic compounds.

[1016] Specific examples of peptidomimetics derived from the polypeptides described in the previous section are presented below. These examples are illustrative and not limiting in terms of the other or additional modifications.

[1017] XXI.E.1. Peptides with a Reduced Isostere Pseudopeptide Bond

[1018] Proteases act on peptide bonds. It therefore follows that substitution of peptide bonds by pseudopeptide bonds

confers resistance to proteolysis. A number of pseudopeptide bonds have been described that in general do not affect polypeptide structure and biological activity. The reduced isostere pseudopeptide bond is a suitable pseudopeptide bond that is known to enhance stability to enzymatic cleavage with no or little loss of biological activity (Couder, et al. (1993), *Int. J. Polypeptide Protein Res.* 41:181-184, incorporated herein by reference). Thus, the amino acid sequences of these compounds may be identical to the sequences of their parent L-amino acid polypeptides, except that one or more of the peptide bonds are replaced by an isostere pseudopeptide bond. Preferably the most N-terminal peptide bond is substituted, since such a substitution would confer resistance to proteolysis by exopeptidases acting on the N-terminus.

[1019] XXI.E.2. Peptides with a Retro-Inverso Pseudopeptide Bond

[1020] To confer resistance to proteolysis, peptide bonds may also be substituted by retro-inverso pseudopeptide bonds (Dalpozzo, et al. (1993), *Int. J. Polypeptide Protein Res.* 41:561-566, incorporated herein by reference). According to this modification, the amino acid sequences of the compounds may be identical to the sequences of their L-amino acid parent polypeptides, except that one or more of the peptide bonds are replaced by a retro-inverso pseudopeptide bond. Preferably the most N-terminal peptide bond is substituted, since such a substitution will confer resistance to proteolysis by exopeptidases acting on the N-terminus.

[1021] XXI.E.3. Peptoid Derivatives

[1022] Peptoid derivatives of polypeptides represent another form of modified polypeptides that retain the important structural determinants for biological activity, yet eliminate the peptide bonds, thereby conferring resistance to proteolysis (Simon, et al., 1992, *Proc. Natl. Acad. Sci. USA*, 89:9367-9371 and incorporated herein by reference). Peptoids are oligomers of N-substituted glycines. A number of N-alkyl groups have been described, each corresponding to the side chain of a natural amino acid.

[1023] XXII. Kits

[1024] The invention provides for diagnostic and therapeutic kits related useful for therapeutic, diagnostic, and research applications. Exemplary kits are disclosed in U.S. Pat. Nos. 5,773,024; 6,017,721; and 6,232,127 B1. The kits of the invention incorporate minicells, and/or include methods of using minicells described herein.

[1025] XXII.A. Diagnostic and Research use Kit Components

[1026] In one embodiment, the invention relates to kits for determining the diagnosis or prognosis of a patient. These kits preferably comprise devices and reagents for measuring one or more marker levels in a test sample from a patient, and instructions for performing the assay. Optionally, the kits may contain one or more means for converting marker level(s) to a prognosis. Such kits preferably contain sufficient reagents to perform one or more such determinations.

[1027] More specifically, a diagnostic kit of the invention comprises any of the following reagents and/or components in any combination.

- [1028] (1) A detectable or detectably labeled first reagent that binds a ligand of interest. The binding reagent can, but need not, be an antibody or an antibody derivative comprising a detectable moiety. The sphingolipid-binding reagent is stored in an openable container in the kit, or is bound to a surface of a substrate such that it is accessible to other reagents. Examples of the latter include test strips.
- [1029] (2) If the first reagent in neither detectable nor detectably labeled, the kit may comprise a detectable or detectably labeled second reagent that binds to the first reagent (e.g., a secondary antibody) or which produces a detectable signal when in close proximity to the first reagent (e.g., as results from fluorescent resonance energy transfer FRET). In either case, the signal produced from the second reagent correlates with the amount of ligand in the sample.
- [1030] (3) One or more positive control reagents. Typically, these reagents comprise a compound that is known to produce a signal in the assay. In one embodiment, the positive control reagents are standards, i.e., comprise a known amount of a detectable or detectably labeled compound, the signal from which may be compared to the signal from a test sample. In addition to serving as positive control reagents, they may be used to develop calibration curves that relate the amount of signal to the known concentration of a detectable or detectably labeled compound. The signal from a test sample is compared to the calibration curve in order to determine what concentration of the detectable or detectably labeled compound corresponds to the signal from the test sample. In this embodiment, the kit provides quantitative measurements of the amount of a ligand in a test sample.
- [1031] (4) One or more negative control reagents. Typically, these control reagents may comprise buffer or another solution that does not contain any of the detectable or detectably labeled first or second reagents and should thus not produce any detectable signal. Any signal that is detected reflects the background level of "noise" in the assay. Another type of negative control reagent contains most of the components necessary for the signal of the assay to be produced, but lacks at least one such component and therefore should not produce a signal. Yet another type of negative control reagent contains all of the components necessary for the signal of the assay to be produced, but also contains an inhibitor of the process that produced the signal.
- [1032] (5) One or more auxiliary reagents for use in the diagnostic assays of the kit, e.g., buffers, alcohols, acid solutions, etc. These reagents are generally available in medical facilities and thus are optional components of the kit. However, these reagents preferably are included in the kit to ensure that reagents of sufficient purity and sterility are used, since the resulting protein conjugates are to be administered to mammals, including humans, for medical purposes, and to provide kits that can be used in situations where medical facilities are not readily available, e.g., when hiking in places located far from medical facilities, or in situations where the presence of these auxiliary reagents allows for the immediate treatment of a patient outside of a medical facility as opposed to treatment that arrives at some later time).
- [1033] (6) Instructions to a person using a kit for its use. The instructions can be present on one or more of the kit components, the kit packaging and/or a kit package insert. XXII.B. Therapeutic Kit Components
- [1034] A therapeutic kit of the invention comprises any of the following reagents and/or components in any combination.
- [1035] (1) One or more therapeutic agents.
- [1036] (2) If the therapeutic agent(s) are not formulated for delivery via the alimentary canal, which includes but is not limited to sublingual delivery, a device capable of delivering the therapeutic agent through some other routes. One type of device for parenteral delivery is a syringe that is used to inject the therapeutic agent into the body of an animal in need of the therapeutic agent. Inhalation devices may also be used.
- [1037] (3) Separate containers, each of which comprises one or more reagents of the kit. In a preferred embodiment, the containers are vials contain sterile, lyophilized formulations of a therapeutic composition that are suitable for reconstitution. Other containers include, but are not limited to, a pouch, tray, box, tube, or the like. Kit components may be packaged and maintained sterilely within the containers.
- [1038] (4) Instructions to a person using a kit for its use. The instructions can be present on one or more of the kit components, the kit packaging and/or a kit package insert. Such instructions include, by way of non-limiting example, instructions for use of the kit and its reagents, for reconstituting lyophilized reagents or otherwise preparing reagents.
- [1039] A preferred kit of the present invention comprises the elements useful for performing an immunoassay. A kit of the present invention can comprise one or more experimental samples (i.e., formulations of the present invention) and one or more control samples bound to at least one pre-packed dipstick or ELISA plate, and the necessary means for detecting immunocomplex formation (e.g., labelled secondary antibodies or other binding compounds and any necessary solutions needed to resolve such labels, as described in detail above) between antibodies contained in the bodily fluid of the animal being tested and the proteins bound to the dipstick or ELISA plate. It is within the scope of the invention that the kit can comprise simply a formulation of the present invention and that the detecting means can be provided in another way.
- [1040] An alternative preferred kit of the present invention comprises elements useful for performing a skin test. A kit of the present invention can comprise at least one pre-packed syringe and needle apparatus containing one or more experimental samples and/or one or more control samples. A kit according to the invention may be designed for both

diagnostic and therapeutic applications. Any combination of the above elements XX.A.(1)-(6) and XX.B.(1)-(4) may be used in a kit, optionally with additional reagents, standards, sample containers, and the like.

[1041] XXIII. Immunogenic Minicells

[1042] XXIII.A. In General

[1043] Minicells are used to immunize subjects. An organism is said to be "immunized" when, after contact with an immunogen, the organism produces antibodies directed to the immunogen, or has increased proliferation or activity of cytotoxic and/or helper T cells, or both. Increased proliferation or activity of T cells may be particularly desirable in the case of parasites that cause a decrease in T cell proliferation.

[1044] The use of minicells to present antigens has several potential advantages. An intact membrane protein can be presented in its native form on the surface of an immunogenic minicell, rather than as a denatured protein or as oligopeptides derived from the amino acid sequence of a membrane protein, which allows for antibodies to be developed that are directed to epitopes which, due to protein folding, occur only in the native protein. The minicell surface may naturally be, or may be modified to be, an adjuvant. Moreover, pharmacokinetic properties of minicells, as discussed elsewhere herein, may be improved relative to other forms of administration.

[1045] The applications of immunogenic minicells include, but are not limited to, research, prophylactic, diagnostic and therapeutic applications.

[1046] In research applications, immunogenic minicells are used to generate antibodies to an antigen displayed on a minicell. Such antibodies are used to detect an antigen, which may be a chemical moiety, molecule, virus, organelle, cell, tissue, organ, or organism that one wishes to study. Classically, such antibodies have been prepared by immunizing an animal, often a rat or a rabbit, and collecting antisera therefrom. Molecular biology techniques can be used to prepare antibodies and antibody fragments, as is described elsewhere herein. Single-chain antibody fragments (scFv) may also be identified, purified, and characterized using minicells displaying a membrane protein or membrane bound chimeric soluble protein.

[1047] In prophylactic applications, immunogenic minicells are used to stimulate a subject to produce antibodies and/or activate T cells, so that the subject is "pre-immunized" before contact with a pathogen or hyperproliferative cell. Thus, in the case of a pathogen, the subject is protected by antibodies and/or T cells that are specifically directed to the pathogen before infection.

[1048] In therapeutic applications, immunogenic minicells are used in immunotherapy.

[1049] Certain aspects of the invention involve active immunotherapy, in which treatment relies on the *in vivo* stimulation of the endogenous host immune system to react against pathogens or tumors due to the administration of agents that cause, enhance or modulate an immune response. Such agents include, but are not limited to, immunogens, adjuvants, cytokines and chemokines.

[1050] Other therapeutic applications involve passive immunotherapy, in which treatment involves the delivery of

agents (such as antibodies or effector cells) that are specifically directed to an immunogen of a pathogen or a hyperproliferative cell, and does not necessarily depend on an intact host immune system. Examples of effector cells include T cells; T lymphocytes, such as CD8+cytotoxic T lymphocytes and CD4+ T-helper tumor-infiltrating lymphocytes; killer cells, such as Natural Killer (NK) cells and lymphokine-activated killer cells.

[1051] XXIII.B. Hyperproliferative Disorders

[1052] The immunogenic minicells of the invention can be used to treat hyperproliferative disorders by inducing an immune response to an antigen associated therewith. The term "hyperproliferative disorder" refers to disorders characterized by an abnormal or pathological proliferation of cells, for example, cancer, psoriasis, hyperplasia and the like.

[1053] For reviews of immunotherapy as applied to hyperproliferative disorders, see Armstrong et al., *Cellular immunotherapy for cancer*, *BMJ* 323:1289-1293, 2001; Evans, *Vaccine therapy for cancer—fact or fiction?*, *Proc R Cell Physicians Edinb* 31:9-16, 2001; Ravindranath and Morton, "Active Specific Immunotherapy with Vaccines," Chapter 61 in: *Holland-Frei Cancer Medicine*, Fifth Edition, Bast, Robert C., et al., editors, B. C. Decker, Inc., Hamilton, 2000, pages 800-814.

[1054] Types of cancers include without limitation fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, chordoma, angiosarcoma, endotheliosarcoma, lymphangiosarcoma, lymphangioendotheliosarcoma, synovioma, mesothelioma, Ewing's tumor, leiomyosarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary adenocarcinomas, cystadenocarcinoma, medullary carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma, choriocarcinoma, seminoma, embryonal carcinoma, Wilms' tumor, cervical cancer, testicular tumor, lung carcinoma, small cell lung carcinoma, bladder carcinoma, epithelial carcinoma, glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, pinealoma, hemangioblastoma, acoustic neuroma, oligodendroglioma, meningioma, melanoma, neuroblastoma, retinoblastoma, leukemia, lymphoma, multiple myeloma, Waldenstrom's macroglobulinemia, and heavy chain disease.

[1055] Tumor specific antigens (TSAs), tumor-associated differentiation antigens (TADAs) and other antigens associated with cancers and other hyperproliferative disorders include, but are not limited to, C1 IAC, a human cancer associated protein (Osther, U.S. Pat. No. 4,132,769); the CA 125 antigen, an antigen associated with cystadenocarcinoma of the ovary, (Hanisch et al., *Carbohydr. Res.* 178:29-47, 1988; O'Brien, U.S. Pat. No. 4,921,790); CEA, an antigen present on many adenocarcinomas (Horig et al., *Strategies for cancer therapy using carcinoembryonic antigen vaccines*, *Expert Reviews in Molecular Medicine*, <http://www-erm-m.cbuc.cam.ac.uk>: 1, 2000); CORA (carcinoma or orosomucoid-related antigen) described by Toth et al. (U.S. Pat. No. 4,914,021); DF3 antigen from human breast carcinoma (Kufe, in U.S. Pat. Nos. 4,963,484 and 5,053,489); DUPAN-2, a pancreatic carcinoma antigen (Lan et al., *Cancer*

Res. 45:305-310, 1985); HCA, a human carcinoma antigen (Cordington et al., U.S. Pat. No. 5,693,763); Her2, a breast cancer antigen (Fendly et al., The Extracellular Domain of HER2/neu Is a Potential Immunogen for Active Specific Immunotherapy of Breast Cancer, Journal of Biological Response Modifiers 9:449-455, 1990); MSA, a breast carcinoma glycoprotein (Tjandra et al., Br. J. Surg. 75:811-817, 1988); MFGM, a breast carcinoma antigen (Ishida et al., Tumor Biol. 10:12-22, 1989); PSA, prostate specific antigen (Nadji et al., Prostatic-specific-antigen, Cancer 48:1229-1232, 1981); STEAP (six transmembrane epithelial antigens of the prostate) proteins (Afar et al., U.S. Pat. No. 6,329,503); TAG-72, a breast carcinoma glycoprotein (Kjeldsen et al., Cancer Res. 48:2214-2220, 1988); YH206, a lung carcinoma antigen (Hinoda et al., Cancer J. 42:653-658, 1988); the p97 antigen of human melanoma (Estin et al., Recombinant Vaccinia Virus Vaccine Against the Human Melanoma Antigen p97 for Use in Immunotherapy, Proc. Natl. Acad. Sci. USA, 85:1052-1056, 1988); and the melanoma specific antigen described by Pfreundschuh in U.S. Pat. No. 6,025,191);

#### [1056] XXIII.B. Intracellular Pathogens

[1057] In certain aspects of the invention, vaccines comprising immunogenic minicells are used to prevent or treat diseases caused by intracellular pathogens. Vaccines may be prepared that stimulate cytotoxic T cell responses against cells infected with viruses including, but not limited to, hepatitis type A, hepatitis type B, hepatitis type C, influenza, varicella, adenovirus, herpes simplex type I (HSV-I), herpes simplex type II (HSV-II), rinderpest, rhinovirus, echovirus, rotavirus, respiratory syncytial virus, papilloma virus, papova virus, cytomegalovirus, echinovirus, arbovirus, huntavirus, coxsackie virus, mumps virus, measles virus, rubella virus, polio virus, human immunodeficiency virus type I (HIV-I), and human immunodeficiency virus type II (HIV-II). Vaccines also may be prepared that stimulate cytotoxic T cell responses against cells infected with intracellular obligates, including but not limited to Chlamydia, Mycobacteria and Rickettsia. Vaccines also may be prepared that stimulate cytotoxic T cell responses against cells infected with intracellular protozoa, including, but not limited to, leishmania, kokzidioa, and trypanosoma.

[1058] The causative agent of Lyme disease, the spirochete *Borrelia burgdorferi*, is also of interest. The outer surface proteins (Osp) A, B and C of *B. burgdorferi* are known antigens that are lipoproteins that associate with membranes. Amino-terminal cysteine residues in Osp proteins are the sites of triacyl lipid modifications that serve as membrane-anchoring moieties. The N-terminal portions of the Osp proteins are highly conserved and are preferred portions for display on immunogenic minicells.

#### [1059] XXIII.C. Eukaryotic Pathogens

[1060] In addition to intracellular pathogens, other eukaryotic pathogens exist and may also be treated using immunogenic minicells displayed antigens therefrom. A number of antigens have been used to develop anti-parasitic vaccines, e.g. the recombinant 45w protein of *Taenia ovis*; EG95 oncosphere proteins of *Echinococcus granulosus*; cathepsin L antigen of the liver fluke, *Fasciola hepatica*; and the H11 antigen of *Haemonchus contortus* (Dalton et al., Parasite vaccines—a reality?, Vet Parasitol 98:149-167, 2001). Other eukaryotic pathogens include, but are not limited to:

[1061] Protozoans, including but not limited to, *Entamoeba histolytica*, a pathogenic amoeba that causes amoebic dysentery and occasionally digests its way through the intestinal wall to invades other organs, which may cause morbidity; *Balantidium coli*, a ciliate that causes diarrhea in humans; *Giardia lamblia*, a flagellate that causes diarrhea and abdominal pain, along with a chronic fatigue syndrome that is otherwise asymptomatic and difficult to diagnose; *Trypanosoma brucei*, a hemoflagellate causing sleeping sickness; and *Trypanosoma cruzi*, the cause of Chagas disease);

[1062] Plasmodia, sporozoan obligate intracellular parasites of liver and red blood cells, including but not limited to *P. falciparum*, the causative agent of malaria. Dozens of *P. falciparum* antigens have been identified, e.g., CSP-1, STARP, SALSA, SSP-2, LSA-1, EXP-1, LSA-3, RAP-1, RAP-2, SERA-1, MSP-1, MSP-2, MSP-3, MSP-4, MSP-5, AMA-1, EBA-175, RESA, GLURP, EMP-1, Pfs25, Pfg27, Pf35, Pf55, Pfs230, Pfg27, Pfs16, Pfs28 and Pfs45/48.

[1063] Helminthes including but not limited to *Ascaris lumbricoides* (roundworm); *Enterobius vermicularis* (pinworm); *Trichuris trichiuria* (whipworm); and *Fasciola hepatica* (liver fluke);

[1064] *Taenia* sp. (tapeworms and cestodes);

[1065] *Schistosoma* (trematodes), such as *Schistosoma mansoni*, which comprises the Sm32 antigen (asparaginyl endopeptidase), which can induce antibody formation in mice (Chlichlia et al., DNA vaccination with asparaginyl endopeptidase (Sm32) from the parasite *Schistosoma mansoni*: anti-fecundity effect induced in mice, Vaccine 20:439-447, 2001); and acetylcholinesterase (Arnon et al., Acetylcholinesterase of *Schistosoma mansoni*-Functional correlates, Protein Science 8:2553-2561, 1999); and

[1066] Ticks and other invertebrates, including but not limited to insects, arachnids, etc. For example, a description of a vaccine against the cattle tick *Boophilus microplus* has been described (Valle et al., The evaluation of yeast derivatives as adjuvants for the immune response to the Bm86 antigen in cattle, BMC Biotechnol. 1:2, 2001)

[1067] XXIII.D. Formulation and Administration of Immunogenic Minicells

[1068] Vaccine formulations of immunogenic minicells include a suitable carrier. Because minicells may be destroyed by digestion, or prevented from acting due to antibody secretion in the gut, they are preferably administered parenterally, including, for example, administration that is subcutaneous, intramuscular, intravenous, or intradermal. Formulations suitable for parenteral administration include aqueous and non-aqueous sterile injection solutions which may contain antioxidants, buffers, and solutes which render the formulation isotonic with the bodily fluid, preferably the blood, of the individual; and aqueous and non-aqueous sterile suspensions which may include suspending agents or thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example, sealed ampules and vials and may be stored in a freeze-dried condition requiring only the addition of the sterile liquid carrier immediately prior to use. The vaccine formulation may also include adjuvant systems for enhancing the immunogenicity of the formulation. Adjuvants are substances that can be used to augment a specific immune response. Nor-

mally, the adjuvant and the composition are mixed prior to presentation to the immune system, or presented separately, but into the same site of the mammal being immunized. Examples of materials suitable for use in vaccine compositions are provided in Osol, A., ed., Remington's Pharmaceutical Sciences, Mack Publishing Co, Easton, Pa. (1980), pp. 1324-1341, which reference is entirely incorporated herein by reference.

[1069] Compositions comprising immunogenic minicells are injected into a human or animal at a dosage of 1-1000 ug per kg body weight. Antibody titers against growth factor are determined by ELISA, using the recombinant protein and horseradish peroxidase-conjugated goat anti-human or animal immunoglobulins or other serologic techniques (e.g., sandwich ELISA). Booster injections are administered as needed to achieve the desired levels of protective antibodies and/or T cells.

[1070] Routes and frequency of administration, as well as dosage, will vary from individual to individual. Between 1 and 10 doses may be administered for a 52-week period. Preferably, 6 doses are administered, at intervals of 1 month, and booster vaccinations may be given periodically thereafter. Alternate protocols may be appropriate for individual patients. In immunotherapy of hyperproliferative disorders, a suitable dose is an amount of a compound that, when administered as described above, is capable of promoting an anti-tumor immune response. Such response can be monitored by measuring the anti-tumor antibodies in a patient or by vaccine-dependent generation of cytolytic effector cells capable of killing the patient's tumor cells in vitro. Such vaccines should also be capable of causing an immune response that leads to an improved clinical outcome (e.g., more frequent remissions, complete or partial or longer disease-free survival) in vaccinated patients as compared to non-vaccinated patients.

[1071] The vaccine according to the invention may contain a single species of immunogenic minicells according to the invention or a variety of immunogenic minicells, each of which displays a different immunogen. Additionally or alternatively, immunogenic minicells may each display and/or express more than one immunogen.

[1072] The summary of the invention described above is non-limiting and other features and advantages of the invention will be apparent from the following detailed description of the invention, and from the claims.

## EXAMPLES

### Example 1

#### Creation of a Minicell-Producing Bacterial Cell Line (MC-T7) that Expresses an Exogenous RNA Polymerase

[1073] In order to maximize the amount of RNA transcription from episomal elements in minicells, a minicell-producing cell line that expresses an RNA polymerase specific for certain episomal expression elements was created. This *E. coli* strain, designated MC-T7, was created as follows.

[1074] The P678-54 *E. coli* strain contains mutations that influence cell division and induce the production of minicells (Adler et al., Proc. Natl. Acad. Sci. 57:321-326 (1967),

Allen et al., Biochem. Biophys. Res. Communi. 47:1074-1079 (1972), Hollenberg et al., Gene 1:33-47 (1976)). The P678-54 strain is resistant to Lambda phage due to a mutation in the *malT* gene (Gottesman, Bacteriophage Lambda: The Untold Story. J. Mol. Biol. 293:177-180, 1999; Friedman, Interactions Between Bacteriophage Lambda and its *Escherichia Coli* Host. Curr. Opin. Genet. Dev. 2:727-738, 1992). Thus, as an initial step, the P678-54 strain was altered so as to be sensitive to Lambda phage so that it could form lysogens of Lambda-DE3 (see below). Wildtype *MalT*-encoding sequences were restored via a HFR (high frequency recombination) conjugation protocol using the G43 *E. coli* strain (CGSC stain 4928).

[1075] Recipient (P678-54) and donor (G43:BW6169) strains were grown overnight in 10 mL of LB media (10 g NaCl, 10 g select peptone 140, and 5 g yeast extract in one liter ddH<sub>2</sub>O). The samples were centrifuged and then concentrated in about 0.2 mL of LB media. The concentrated samples were combined and incubated with slow rotation for 30 minutes at 30° C., and were then plated on LB agar plates that contained streptomycin (50 µg/mL) and tetracycline (50 µg/mL). (Ampicillin, streptomycin, tetracycline, and all other chemicals were purchased from Sigma Chemical (St. Louis, Mo.) unless otherwise indicated.) Recipient cells were resistant to streptomycin and donor cells were resistant to tetracycline; only conjugates, which contained both resistance genes, were able to grow on the LB agar plates that contained streptomycin (50 µg/mL) and tetracycline (50 µg/mL).

[1076] Putative conjugates were screened for Lambda phage sensitivity using a cross streak technique, in which putative colonies were cross-streaked on an LB agarose plate (streptomycin, 50 µg/mL, and tetracycline, 50 µg/mL) that had been streaked with live Lambda phage. The streaked conjugate colonies were streaked perpendicular to the Lambda phage streak; if a conjugate was sensitive to Lambda phage infection then, upon contact with the Lambda phage streak, there was cell lysis and thus less or no bacterial growth. Thus, in the case of conjugates that were sensitive to Lambda phage, there was decreased bacterial growth "downstreak" from the phage streak.

[1077] The conjugate *E. coli* that were found to be sensitive to Lambda phage infection were then used to create Lambda lysogens. Lysogenization is a process during which Lambda phage incorporates its genome, including exogenous genes added thereto, into a specific site on the chromosome of its *E. coli* host cell.

[1078] The DE3 gene, which is present in the genome of the Lambda phage used to create lysogens, encodes RNA polymerase from bacteriophage T7. Lysogenation was carried out using the DE3-Lysogenation kit (Novagen, Madison, Wis.) essentially according to the manufacturer's instructions. A T7 polymerase dependent tester phage was used to confirm the presence and expression of the DE3 gene on the bacterial chromosome. The T7-dependent tester phage can only form plaques on a bacterial known in the presence of T7 polymerase. The phage uses a T7 promoter for expression of its essential genes. Therefore in a plaque-forming assay only cells which express T7 polymerase can be lysed by the tester phage and only these cells will allow for the formation of plaques. As is described in more detail herein, episomal expression elements that are used in mini-

cells may be designed such that transcription and translation of a cloned gene is driven by T7 RNA polymerase by utilizing expression sequences specific for the T7 RNA polymerase.

### Example 2

#### Cloning of Rat Edg-1 into the pCAL-c Expression Vector

##### [1079] Materials

[1080] Taq Polymerase, PCR Buffers, and PCR reagents were purchased from Roche Molecular Biochemicals (Indianapolis, Ind.). All restriction enzymes were purchased from Gibco BRL (Grand Island, N.Y.) and Stratagene (La Jolla, Calif.). QIAprep mini and maxi kits, PCR purification Kits, RNeasy miniprep kits, and the One Step RT-PCR Kit were purchased from QIAGEN (Valencia, Calif.). The GeneClean Kit was purchased from BIO 101 (Carlsbad, Calif.). IPTG (isopropyl-beta-D-thiogalactopyranoside), T4 DNA Ligase, LB Media components and agarose were purchased from Gibco BRL. The pCAL-c prokaryote expression vector and competent cells were purchased from Stratagene.

[1081] The pCAL-c expression vector has a structure in which an ORF may be operably linked to a high-level (but T7 RNA polymerase dependent) promoter, sequences that bind the *E. coli* Lac repressor, and the strong T7 gene 10 ribosome-binding site (RBS). The *LacI* repressor is also encoded by an expressed from the pCAL-c vector. As long as it is bound to its recognition sequences in the pCAL-c

expression element, the *lac* repressor blocks transcription from the T7 promoter. When an inducing agent, such as IPTG is added, the *lac* repressor is released from its binding sites and transcription proceeds from the T7 promoter, provided the T7 RNA polymerase is present. After induction, the cloned and expressed protein may constitute the majority of newly expressed cellular proteins due to the efficient transcription and translation processes of the system.

##### [1082] Amplification

[1083] The first step in cloning rat Edg-1 (rEDG-1) into an expression vector was to design primers for amplification via PCR (polymerase chain reaction). PCR primers were designed using the rat Edg-1 sequence (Nakajima et al., *Biophys. J.* 78:319A, 2000) in such a manner that they contained either sites for *NheI* (GCTAGC) or *BamHI* (GGATCC) on their five prime ends. The upstream primer had the sequence of SEQ ID NO:31. The three prime downstream primer (SEQ ID NO:32) also contained a stop codon, as the pCAL-c vector contains a Calmodulin Binding Protein (CBP) "tag" at its carboxyl terminus which was not intended to be incorporated into the rat Edg-1 polypeptide in this expression construct. The primer and resulting PCR products were designed so that the five prime end of the rat Edg-1 ORF was in frame with the methionine start codon found in the pCAL-c vector.

##### [1084] Oligonucleotide Primer Sequences for Cloning into pCAL-c:

###### Edg1/pCAL-c construct primers:

###### Upstream primer

5'-AATTGCTAGCTCCACCAGCATCCCAGTGGTTA-3' (SEQ ID NO:31)

###### Downstream primer

5'-AATTGGATCCTTAAGAAGAAGAAATTGACGTTT-3' (SEQ ID NO:32)

###### Edg1/CBP fusion construct primers:

###### Upstream primer

5'-AATTGCTAGCTCCACCAGCATCCCAGTGGTTA-3' (SEQ ID NO:31)

###### Downstream primer

5'-AATTGGATCCAGAAGAAGAAATTGACGTTTCCA-3' (SEQ ID NO:33)

###### Edg1/His6 construct primers:

###### Upstream primer

5'-AATTGCTAGCTCCACCAGCATCCCAGTGGTTA-3' (SEQ ID NO:31)

###### Downstream primer

5'-AATTGGATCCTTAATGATGATGATGATGATGATGAGAAGAAGAAATTGACGTTTCC-3' (SEQ ID NO:34)

###### Edg3/rtPCR primers:

###### Upstream primer

5'-TTATGGCAACCACGCACGCAGG-3' (SEQ ID NO:35)

###### Downstream primer

5'-AGACCGTCACTTGACAGGAC-3' (SEQ ID NO:36)

###### Edg3/pCAL-c construct primers:

###### Upstream primer

5'-AATTGCTAGCACGCACGCGCAGGGCCACCCGC-3' (SEQ ID NO:37)

###### Downstream primer

**-continued**

5'-AATTGGTACCTCACTTGCGAGGACCCATTCTG-3' (SEQ ID NO:38)

**Edg3/His6 construct primers:****Upstream primer**

5'-AATTGCTAGCACGCACGCGCAGGGGCACCCGC-3' (SEQ ID NO:39)

**Downstream primer**

5'-AATTGGTACCTCAATGATGATGATGATGATGCTTGCGAGGACCCCTTCTG-3' (SEQ ID NO: 16)

**GFP/pCAL-c construct primers:****Upstream primer**

5'-GGTCGCCACCATGGTGAGCAA-3' (SEQ ID NO:40)

**Downstream primer**

5'-TTAAGGATCCTTACTTGTACAGCTCGTCCAT-3' (SEQ ID NO:41)

**GFP/CBP construct primers:****Upstream primer**

5'-GGTCGCCACCATGGTGAGCAA-3' (SEQ ID NO:42)

**Downstream primer**

5'-TTAAGGATCCCTTGTACAGCTCGTCCATGCC-3' (SEQ ID NO:43)

**Notes:**

Restriction endonuclease sites are underlined

Stop codons are double underlined

**[1085]** The primers were used to amplify the rEdg-1 DNA ORF using the polymerase chain reaction (PCR). The template used for amplification was mRNA isolated from rat muscle tissue using the RNeasy Miniprep Kit (Qiagen) and was carried out essentially according to the manufacturer's protocol. Both the rtPCR and PCR amplification steps were carried out in a single reaction using the One Step RT-PCR Kit (Qiagen). The resulting rat Edg-1 PCR fragment was purified using the PCR Purification Kit (Qiagen). The amplified double stranded rEdg-1 DNA sequence contained the NheI site at the 5-prime end and the BamHI site at the 3-prime end. This amplified rEdg-1 fragment was used for cloning into the pCAL-c expression vector.

**[1086]** The pCAL-c expression vector contains NcoI, NheI, and BamHI restriction sites in its multiple cloning site. In order to insert rEdg-1-encoding sequence into the expression vector, the rEdg-1 PCR fragment and the pCAL-c expression vector were digested with NheI and BamHI restriction enzymes for one hour at 37° C. The reaction mixture for the digestion step consisted of 1 µg of DNA, 1× restriction buffer, and 1 µL of each enzyme. The reaction mixture was brought to a final volume of 20 µL with ddH<sub>2</sub>O (dd, double distilled). After 45 minutes, 1 µL of Calf Intestine Alkaline Phosphatase (CIAP) was added to the pCAL-c reaction mixture in order to remove the terminal phosphates from the digested plasmid DNA. The reactions were incubated for an additional 15 minutes at 37° C. The digested DNA samples were then run on a 1% TAE (Tris-acetate/EDTA electrophoresis buffer) agarose gel at 130 volts for 45 minutes. The bands were visualized with UV light after the gel was stained with ethidium bromide.

**[1087]** The appropriate bands were cut out of the gel for purification using the GeneClean Kit (BIO101). The Purified DNA fragments were then quantified on a 1% TAE agarose gel. For the ligation reaction, ratios of insert to vector of 6:1 and 3:1 were used. A negative control comprising vector only was also included in the ligation reactions. The reaction mixtures contained insert and vector DNA, 4 µL Ligase buffer, and 2 µL Ligase. The reaction was brought up to a

final volume of 20 µL with ddH<sub>2</sub>O. The ligation was carried out at room temperature for about 2 hours. Ten (10) µL of the ligation reaction mixture was used for subsequent transformation steps.

**[1088]** Ligated DNA was introduced into Epicurian Coli XL1-Blue competent cells using the heat shock transformation technique as follows. The ligation mixture was added to 100 µL of competent cells, placed on ice, and was incubated for about 30 minutes. The cells were then heat shocked at 37° C. for 1 minute and put back on ice for 2 minutes. Following heat shock, 950 µL of room temperature LB media was added to the cells and the cells were shaken at 37° C. for 1 hour. Following the 1-hour agitation the cells were pelleted for one minute at 12000 rpm in a Eppendorf 5417C microcentrifuge. The supernatant was carefully poured off so that about 200 µL remained. The cells were then resuspended in the remaining LB media and spread on 100×15 mm LB agarose plates containing 50 µg/mL ampicillin. The plates were incubated overnight at 37° C. Colonies were counted the following day, and the ratio of colonies between the negative control and the ligated samples was determined. A high ratio of the number of colonies when the ligation mixture was used to transform cells, as contrasted to the number of negative control colonies indicated that the cloning was successful. Transformed colonies were identified, isolated, and grown overnight in LB media in the presence of ampicillin. The resulting bacterial populations were screened for the presence of the Edg-1-pCAL-c expression construct.

**[1089]** Plasmid DNA was isolated from the cells using the QIAprep Spin Miniprep Kit (Qiagen). Isolated Edg-1-pCAL-c constructs were screened using the restriction enzyme ApaI, which digests the Edg-1-pCAL-c construct at two different sites: one in the Edg-1 coding sequence and one in the pCAL-c vector itself. The plasmid preparations were digested with ApaI electrophoresed on a 1% TAE agarose gel and visualized using uv light and ethidium bromide staining. The predicted sizes of the expected DNA fragments were 2065 bp and 4913 bp. As shown in **FIG. 3**,

bands of the predicted size were present on the gel. The entire Edg-1-pCAL-c construct was sequenced in order to confirm its structure. This expression construct, a pCAL-c derivative that contains the rat Edg-1 ORF operably linked to a T7 promoter and lac repressor binding sites, is designated "prEDG-1" herein.

#### Example 3

##### Construction of Rat Edg-1-CBP Fusion Protein

[1090] In order to detect rat Edg-1 protein expression, rEdg-1 coding sequences were cloned into the pCAL-c vector in frame with a CBP fusion tag. The cloning strategy for the rEdg-1-CBP construct was performed essentially as described for the Edg-1-pCAL-c construct with the following differences. The PCR primers (SEQ ID NOS:3 and 5) were as described for the Edg-1-pCAL-c cloning except for the omission of the stop codon in the downstream primer (SEQ ID NO:33). The removal of the stop codon is required for the construction of the Edg-1-CBP fusion protein. The pCAL-c vector is designed so that, when the BamHI site is used for insertional cloning, and no stop codon is present in an ORF inserted into the pCAL-c expression vector the cloned ORF will be in-frame with the CBP fusion tag. Because the three prime downstream primer did not contain a stop codon, a CBP fusion tag could be cloned in-frame with the Edg-1 ORF. Other cloning steps were performed essentially as described before. The resulting plasmid, a pCAL-c derivative that comprises an ORF encoding a rat Edg-1-CBP fusion protein operably linked to a T7 promoter and lac repressor binding sites, is designated "prEDG-1-CBP" herein.

#### Example 4

##### Cloning of a His-Tagged Rat Edg-1 into pCAL-c Expression Vector

[1091] The rEdg-1 protein was manipulated to generate a fusion protein having a 6× His tag at its carboxyl terminus. A "6× His tag" or "His tag" is an amino acid sequence consisting of six contiguous histidine residues that can be used as an epitope for the binding of anti-6× His antibodies, or as ligand for binding nickel atoms. The His-tagged rEdg-1 fusion protein is used to detect rEdg-1 protein expression in the minicell expression system environment.

[1092] The rEdg-1-6× His construct was cloned using the strategy described above for the construction of the rEdg-1-pCAL-c expression construct (prEDG-1), with the upstream primer having the sequence of SEQ ID NO:3, but with the exception that the three prime downstream primer (SEQ ID NO:34) was designed to contain six histidine codons followed by a stop codon. The 18 base pair 6× His tag was incorporated into the carboxyl terminus of the Edg-1 protein as expressed from the pCAL-c vector. Subsequent cloning procedures (PCR, restriction digest, gel purification, ligation, transformation, etc.) were performed as described previously for the Edg-1-pCAL-c construct (prEDG-1). The resulting plasmid, a pCAL-c derivative that comprises an ORF encoding a carboxy-terminal His-tagged rat Edg-1-CBP fusion protein operably linked to a T7 promoter and lac repressor binding sites, is designated "prEDG-1-6× His" herein.

#### Example 5

##### Amplification and Cloning of Rat Edg-3 Sequences

[1093] The Edg-3 full length coding sequence was amplified via PCR from rat skeletal muscle mRNA using primers (SEQ ID NOS:35 and 36) designed from the known mouse sequence (Genbank accession NM\_010101). The mRNA used as a template for the amplification reaction was isolated using the RNeasy Miniprep Kit (Qiagen). Both the rtPCR and PCR amplification steps were carried out in a single reaction using the One Step RT-PCR Kit (Qiagen). The rEdg-3 PCR products were visualized with UV after electrophoresis in 1% TAE agarose gels and ethidium bromide staining.

[1094] The predicted size of the amplified PCR products is 1145 base pairs. An appropriately-sized DNA band was isolated from the TAE gel and purified using the GeneClean Kit (BIO101). The purified band was ligated to the pCR3.1 vector using the TA-cloning kit (Invitrogen). Other cloning steps were carried out as described previously for the cloning of the rEdg-1-pCAL-c construct (prEDG-1) with the exception that the samples were screened using the EcoRI restriction enzyme. The expected sizes of the digested bands were 1145 base pairs and 5060 base pairs. Positive clones were analyzed by automated sequencing. The nucleotide sequences were analyzed using BLAST searches from the NCBI web site ([www.ncbi.nlm.nih.gov/](http://www.ncbi.nlm.nih.gov/)). The predicted full length rat Edg-3 amino acid sequence was assembled from the nucleotide sequencing data using *in silico* translation. The pCR3.1 vector comprising the rat Edg-3 ORF is designated "pCR-rEDG-3" herein.

#### Example 6

##### Cloning of Rat Edg-3 Coding Sequences into the pCAL-c Expression Vector

[1095] In order to express it in the minicell expression system, the rat Edg-3 ORF was cloned into the pCAL-c expression vector. The cloning strategy used was as described above for the cloning of the rat Edg-1 gene into the pCAL-c vector with the following exceptions. The primers used for PCR amplification were designed from the rat Edg-3 sequence and contained sites for the restriction enzymes NheI and KpnI (GGTACC). The NheI site was added to the five prime upstream primer (SEQ ID NO:37) and the KpnI site was added to the three prime downstream primer; SEQ ID NO:38). The NheI and KpnI restriction enzymes were used for the digestion reaction. The reaction mixture for the digestion step consisted of 1 μg of DNA, 1× restriction buffer (provided with the enzyme), and 1 μL of each enzyme. Plasmid preparations were screened by digestion with NheI and KpnI. The digested plasmid DNA was electrophoresed on a TAE agarose gel and visualized by UV after staining with ethidium bromide. The resultant band sizes were predicted to be 1145 base pairs and 5782 base pairs. The positive plasmid clones were analyzed with automated sequencing. The resulting plasmid, a pCAL-c derivative that comprises an ORF encoding a rat Edg-3 protein operably linked to a T7 promoter and lac repressor binding sites, is designated "pEDG-3" herein.



## Example 7

## Cloning of a His-Tagged Rat Edg-3 into the pCAL-c Expression Vector

[1096] In order to detect expression of the rat Edg-3 protein in the minicell expression system, the rat Edg-3 coding sequence was manipulated so as to contain a 6× His tag at the carboxyl terminus of the protein. The cloning strategy used to create this construct was essentially the same as described above for the rEdg-3-pCAL-c (prEDG-3) construct cloning, with the upstream primer having the sequence of SEQ ID NO:37, with the exception that the three-prime downstream primer (SEQ ID NO: 18) was designed to contain a 6× His coding sequence followed by a stop codon, which allowed for the incorporation of the 6× His amino acid sequence onto the carboxyl terminus of the Edg-3 receptor protein. Other cloning and screening steps were performed as described above. The resulting plasmid, a pCAL-c derivative that comprises an ORF encoding a carboxy-terminal His-tagged rat Edg-3 fusion protein operably linked to a T7 promoter and lac repressor binding sites, is designated “prEDG-3-6xHis” herein.

## Example 8

## GFP Cloning into pCAL-c Expression Construct

[1097] Cloning of GFP-encoding nucleotide sequences into the pCAL-c vector was performed in order to produce an expression construct having a reporter gene that can be used to detect protein expression (GFP, green fluorescent protein). The cloning strategy used was essentially the same as the cloning strategy described above with the following exceptions. The template used for PCR amplification was the peGFP plasmid “construct” (GFP construct sold by Clontech). The primers used for amplification were designed from the GFP coding sequence and contained sites for the restriction enzymes NcoI and BamHI. The NcoI site was added to the five prime upstream primer (SEQ ID NO:40) and the BamHI site was added to the three prime downstream primer; see SEQ ID NO:41) The NcoI and BamHI restriction enzymes were used for the digestion reaction. The reaction mixture for the digestion step consisted of 1 μg of DNA, 1× restriction buffer (provided with the enzyme), and 1 μL of each enzyme. The screening of the plasmid preparations was carried out using NcoI and BamHI. Digested plasmid preparations were electrophoresed and visualized on TAE agarose gels with UV after staining with ethidium bromide. Restriction products having the predicted sizes of 797 and 5782 base pairs were seen. Positive plasmid clones were sequenced using an automated sequencer. The resulting plasmid, a pCAL-c derivative that comprises an ORF encoding a rEdg-3-GFP fusion protein operably linked to a T7 promoter and lac repressor binding sites, is designated “prEDG-3-GFP” herein.

## Example 9

## Design Construction of Control Expression Elements

[1098] Control expression elements used to detect and quantify expression of proteins in minicells were proposed. These controls direct the expression of detectable proteins. An expression element used as positive control is pPTC12,

which is supplied with the pCAL-c expression vector from Stratagene. This construct contains an ORF encoding a fusion protein comprising beta-galactosidase linked to CBP. Induction of expression of pTC12 should result in the production of a protein of about 120 kD, and this protein is detected via its enzymatic activity or by using antibodies directed to epitopes on the beta-galactosidase or CBP polypeptide.

[1099] A GFP fusion construct was created and used as a positive control for the CBP detection kit. This construct was a positive control for induction of protein expression in the minicell expression system. The cloning strategy used to create the construct was essentially the same as that used for the cloning of the GFP into the pCAL-c expression vector, with the exception that the three prime downstream primer did not contain a stop codon; this allowed for the in frame incorporation of the CBP fusion tag to the GFP protein. The upstream primer had the sequence of SEQ ID NO:42, and the downstream primer had the sequence SEQ ID NO:43. The nucleotide sequence of the expression element was confirmed using an automated sequencer. The resulting plasmid, a pCAL-c derivative that comprises an ORF encoding GFP operably linked to a T7 promoter and lac repressor binding sites, is designated “pGFP-CBP” herein.

## Example 10

Introduction of pCAL-c Expression Constructs into the MC-T7 *Escherichia coli* Strain

[1100] The MC-T7 *E. coli* strain was made competent using the CaCl<sub>2</sub> technique. In brief, cells were grown in 40 mL LB medium to an OD<sub>600</sub> of 0.6 to 0.8, and then centrifuged at 8000 rpm (7,700 g) for 5 min at 4° C. The pellet was resuspended in 20 mL of cold CaCl<sub>2</sub> and left on ice for five minutes. The cells were then centrifuged at 8000 rpm (7,700 g) for 5 min at 4° C. The cell pellet was resuspended in 1 mL of cold CaCl<sub>2</sub> and incubated on ice for 30 min. Following this incubation 1 mL of 25% glycerol was added to the cells and they were distributed and frozen in 200 μL aliquots. Liquid nitrogen was used to freeze the cells. These cells subsequently then used for the transformation of expression constructs.

## Example 11

## Preparation of Minicells

[1101] To some degree, the preparation of minicells varied according to the type of expression approach that is used. In general, there are two such approaches, although it should be noted from the outset that these approaches are neither limiting nor mutually exclusive. One approach is designed to isolate minicells that already contain an expressed therapeutic protein or nucleic acid. Another approach is designed to isolate minicells that will express the protein or nucleic acid in the minicell following isolation.

[1102] *E. coli* are inoculated into bacterial growth media (e.g., Luria broth) and grown overnight. After this, the overall protocol varies with regards to methods of induction of expression. The minicell producing cultures used to express protein post isolation are diluted and grown to the desired OD<sub>600</sub> or OD<sub>450</sub>, typically in the log growth phase of bacterial cultures. The cultures are then induced with IPTG and then isolated. The IPTG concentration and exposure depended on which construct was being used, but was usually about 500 μM final for a short time, typically about

4 hours. This treatment results in the production of the T7 polymerase, which is under control of the LacUVR5 promoter, which is repressed by the LacI repressor protein. IPTG relieves the LacI repression and thus induces expression from the LacUVR5 promoter which controls expression of the T7 polymerase from the chromosome. This promoter is “leaky” that is, there is always a basal level of T7 polymerase which can be selected for or against so that the induction before isolation is not required. (This induction step is not required if a non-T7 expression system is used, as the reason for this step is to express the T7 RNA polymerase in the minicell-producing cells so that the polymerase and molecules segregate with the minicell.)

[1103] The *E. coli* cultures that produce minicells containing a therapeutic protein or nucleic acid have different induction protocols. The overnight cultures are diluted as described above; however, in the case of proteins that are not toxic to the parent cells, this time the media used for dilution already contains IPTG. The cultures are then grown to mid-log growth and minicells are isolated. These cultures produce the therapeutic protein or nucleic acid as they grow, and the minicells derived therefrom contain the therapeutic protein or nucleic acid.

[1104] Alternatively or additionally, IPTG is added and expression is induced after the isolation of minicells. In the case of non-toxic proteins or nucleic acids that are expressed from expression elements in minicells, this treatment enhances production of the eposimally encoded gene product. In the case of toxic gene products induction post-isolation is preferred.

#### Example 12

##### Minicell Isolation

[1105] Minicells were isolated from the minicell producing MC-T7 strain of *E. coli* using centrifugation techniques. The protocol that was used is essentially that of Jannatipour et al. (Translocation of *Vibrio Harveyi* N,N'-Diacetylchitobiose to the Outer Membrane of *Escherichia Coli*, *J. Bacteriol.* 169: 3785-3791, 1987) and Matsumura et al. (Synthesis of Mot and Che Products of *Escherichia coli* Programmed by Hybrid ColE1 Plasmids in Minicells, *J. Bacteriol.* 132:996-1002, 1977).

[1106] In brief, MC-T7 cells were grown overnight at 37° C. in 2 to 3 mL of LB media containing ampicillin (50 µg/mL), streptomycin (50 µg/mL), and tetracycline (50 µg/mL) (ampicillin was used only when growing MC-T7 cells containing a pCAL-c expression construct). The cells were diluted 1:100 in a total volume of 100 to 200 mL LB media with antibiotics, and grown at 37° C. until they reached an OD<sub>600</sub> of 0.4 to 0.6, which is roughly beginning of the log growth phase for the MC-T7 *E. coli*. During this incubation the remainder of the overnight culture was screened for the presence of the correct expression construct using the techniques described above. When the cultures reached the appropriate OD<sub>600</sub> they were transferred to 250 mL GS3 centrifuge bottles and centrifuged (Beckman centrifuge) at 4500 rpm (3,500 g) for 5 min. At this point the supernatant contains mostly minicells, although a few relatively small whole cells may be present.

[1107] The supernatant was transferred to a clean 250 mL GS3 centrifuge bottle and centrifuged at 8000 rpm (11,300 g) for 10 min. The pellet was resuspended in 2 mL of 1× BSG (10× BSG: 85 g NaCl, 3 g KH<sub>2</sub>PO<sub>4</sub>, 6 g Na<sub>2</sub>HPO<sub>4</sub>, and 1 g gelatin in 1 L ddH<sub>2</sub>O) and layered onto a 32 mL 5 to 20%

continuous sucrose gradient. The sucrose gradient was made with sucrose dissolved in 1× BSG.

[1108] The sucrose gradient was then loaded in a Beckman SW24 rotor and centrifuged in a Beckman Ultracentrifuge at 4500 rpm (9,000 g) for 14 min. Following ultracentrifugation a single diffuse band of minicells was present. The top two thirds of this band was aspirated using a 10 mL pipette and transferred to a 30 mL Oakridge tube containing 10 mL of 1× BSG. The sample was then centrifuged at 13,000 rpm (20,400 g) for 8 min. Following centrifugation, the pellet was resuspended in 2 mL 1× BSG, and the resuspended cells were loaded onto another 5 to 20% sucrose gradient. This sucrose gradient was centrifuged and the minicells were collected as described above. The sucrose gradient procedure was repeated a total of three times.

[1109] Following the final sucrose gradient step the entire minicell band was collected from the sucrose gradient and added to a 30 mL Oakridge tube that contained 10 mL of MMM buffer (200 mL 1× M9 salts, 2 mL 20% glucose, and 2.4 mL DIFCO Methionine Assay Medium). This minicell solution was centrifuged at 13,000 rpm (20,400 g) for 8 min. The pellet was resuspended in 1 mL of MMM Buffer.

[1110] The concentration of minicells was determined using a spectrophotometer. The OD<sub>450</sub> was obtained by reading a sample of minicells that was diluted 1:100.

#### Example 13

##### Other Methods to Prepare and Isolate Minicells

[1111] By way of non-limiting example, induction of *E. coli* parental cells to form minicells may occur by overexpression of the *E. coli* *ftsZ* gene. To accomplish this both plasmid-based and chromosomal overexpression constructs were created that place the *ftsZ* gene under the control of various regulatory elements (Table 6).

TABLE 6

##### REGULATORY CONSTRUCTS CONTROLLING FTSZ EXPRESSION.

Regulatory region	inducer	[inducer]	SEQ ID NO.:
Para::ftsZ	Arabinose	10 mM	1, 3
Prha::ftsZ	Rhamnose	1 mM	2, 4
Ptac::ftsZ	IPTG	30 µM	5, Garrido et al. <sup>a</sup>

<sup>a</sup>Garrido, T. et al. 1993. Transcription of *ftsZ* oscillates during the cell cycle of *Escherichia coli*.

[1112] Oligonucleotide Names and PCR Reactions use the Following Format:

[1113] “gene-1” is N-terminal, 100% homology oligo for chromosomal or cDNA amplification

[1114] “gene-2” is C-terminal, 100% homology oligo for chromosomal or cDNA amplification

[1115] “gene-1-RE site” is same sequence as gene-1 with additional residues for remainder of sequence, RE sites, and/or chimeric fusions.

[1116] “gene-2-RE site” is same sequence as gene-1 with additional residues for remainder of sequence, RE sites, and/or chimeric fusions.

[1117] Use “gene-1, 2” combo for chromosomal/cDNA amplification and “gene-1 RE site, gene-2-RE site” to amplify the mature sequence from the “gene-1, 2” gel-purified product.

TABLE 7

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 6 CONSTRUCTS		
SEQ ID NO.:	Primer name	5' to 3' sequence
44	FtsZ-1	CCAATGGAACCTACCAATGACGCGG
45	FtsZ-2	GCTTGCTTACGCAGGAATGCTGGG
46	FtsZ-1-PstI	CGCGGCTGCAGATGTTGAACCAATGGAACCTACCAATGACGCGG
47	FtsZ-2-XbaI	GCGCCTCTAGATTATTAATCAGCTTGCCACGCAGGAATGCTGGG

[1118] Table 7 oligonucleotide sequences are for use in cloning *ftsZ* into SEQ ID NO.:1 and 2 (insertions of *ftsZ* behind the arabinose promotor (SEQ ID NO.: 1) and the rhamnose promotor (SEQ ID NO.: 2).

TABLE 8

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR FTSZ CHROMOSOMAL DUPLICATION CONSTRUCTS		
SEQ ID NO.:	Primer name	5' to 3' sequence
48	Kan-1	GCTAGACTGGGCGGTTTTATGGACAGCAAGC
49	Kan-2	GCGTTTAATAATTCAGAAGAACTCGTCAAGAAGGCG
50	Kan-1-X-frt	GCGCCTACTGACGTAGTTCGACCGTCGGACTAGCGAAGTTCCTATACTTTCTAGAGAATAGGAACCTTCGCTAGACTGGCGGTTT-TATGGACAG-CAAGC
51	Kan-2-intD-frt	CAAGATGCTTTGCCTTTGTCTGAGTTGATACTGGCTTTGGAAAGTTC-CTATCTCTCTA-GAAAGTATAG-GAACTTCGCGTTAATAATTCA-GAA-GAACTCGT-CAAGAAGGCG
52	AraC-1	CGTTACCAATTATGACAACCTTGACGG
53	RhaR-1	TTAATCTTTCTGCGAATTGAGATGACGCC
54	LacI <sup>q</sup> -1	CGTTACCAATTATGACAACCTTGACGG
55	Ara-1-intD	GCCTGCATTGCGGCGCTTCAGTCTCCGCTGCATACTGTCCCGTTACCAAT-TATGACAACCT-TGACGG
56	RhaR-1-intD	GCCTGCATTGCGGCGCTTCAGTCTCCGCTGCATACTGTCCTTAATCTTTCTGCGAATTGAGATGACGCC
57	LacI <sup>q</sup> -1-intD	GCCTGCATTGCGGCGCTTCAGTCTCCGCTGCATACTGTCCTTAATAAAGT-GAGTCGATAT-TGCTTTTGT-TGACCAG
58	FtsZ-1-X	GCCTGCATTGCGGCGCTTCAGTCTCCGCTGCATACTGTCCCGTTACCAAT-TATGACAACCT-TGACGG

[1119] In like fashion, the *ftsZ* gene was amplified from SEQ ID NO.: 1, 2 and *Ptac::ftsZ* (Garrido, T. et al. 1993. Transcription of *ftsZ* oscillates during the cell cycle of *Escherichia coli*. *EMBO J.* 12:3957-3965) plasmid and chromosomal constructs, respectively using the following oligonucleotides:

[1120] For amplification of *araC* through *ftsZ* of SEQ ID NO.: 1 use oligonucleotides:

[1121] *AraC-1*

[1122] *FtsZ-2*

[1123] For amplification of *rhaR* through *ftsZ* of SEQ ID NO.: 2 use oligonucleotides:

[1124] *RhaR-1*

[1125] *FtsZ-2*

[1126] For amplification of *lacI<sup>q</sup>* through *ftsZ* of *Ptac::ftsZ* (Garrido, T., et al.) use oligonucleotides:

[1127] *lacI<sup>q</sup>-1*

[1128] *ftsZ-2*

[1129] The above amplified DNA regions were gel-purified and used as template for the second round of PCR using oligonucleotides containing homology with the *E. coli* chromosomal gene *intD* and on the other end with random sequence termed "X". Oligonucleotides used in this round of PCR are shown below:

[1130] For amplification of *araC* through *ftsZ* from SEQ ID NO.: 1 to contain homology to *intD* and the random X use oligonucleotides:

[1131] *AraC-1-intD*

[1132] *FtsZ-1-X*

[1133] For amplification of *rhaR* through *ftsZ* from SEQ ID NO.: 2 to contain homology to *intD* and the random X use oligonucleotides:

[1134] *RhaR-1-intD*

[1135] *FtsZ-1-X*

[1136] For amplification of *lacI<sup>q</sup>* through *ftsZ* from *Ptac::ftsZ* to contain homology to *intD* and the random X use oligonucleotides:

[1137] *LacI<sup>q</sup>-1-intD*

[1138] *FtsZ-1-X*

[1139] The PCR products from these PCR reactions are as shown below:

[1140] *intD—araC—Ara promotor—ftsZ—"X"*

[1141] *intD—rhaRS—Rha promotor—ftsZ—"X"*

[1142] *intD—lacI<sup>q</sup>—Ptac promotor—ftsZ—"X"*

[1143] To amplify the mature complexes, the following regions were mixed and amplified with the coupled oligonucleotide sequence primers:

SEQ ID NO.: 3 was produced using:

```

intD-araC-Ara promotor-ftsZ-"X"
AraC-1-intD | "X"-frt-Kan-frt-intD
Kan-2-intD-frt |
↓
intD-araC-Ara promotor-ftsZ-"X"-frt-Kan-frt-intD

```

SEQ ID NO.: 4 was produced using:

```

intD-rhaRS-Rha promotor-ftsZ-"X"
RhaR-1-intD | "X"-frt-Kan-frt-intD
Kan-2-intD-frt |
↓
intD-rhaRS-Ara promotor-ftsZ-"X"-frt-Kan-frt-intD

```

SEQ ID NO.: 5 was produced using:

```

intD-lacIq-Ptac promotor-ftsZ-"X"
lacIq-1-intD | "X"-frt-Kan-frt-intD
Kan-2-intD-frt |
↓
intD-lacIq-Ptac promotor-ftsZ-"X"-frt-Kan-frt-intD

```

[1144] These expression constructs may be expressed from the plasmid, placed in single copy, replacing the native *ftsZ* copy on the *E. coli* chromosome (Garrido, T., et al. 1993. Transcription of *ftsZ* oscillates during the cell cycle of *Escherichia coli*. *EMBO J.* 12:3957-3965), or in duplicate copy retaining the native *ftsZ* copy while inserting one of the expression constructs in Table 6 into the *intD* gene on the same chromosome. Chromosomal duplications were constructed using the RED recombinase system (Katsenko, K. A., and B. L. Wanner. One-Step Inactivation of Chromosomal Genes in *Escherichia coli* K-12 Using PCR Products. *Proc. Natl. Acad. Sci.* 97:6640-6645. 2000) and are shown in SEQ ID NO 3-5. The later constructs allow native replication during non-minicell producing conditions, thus avoiding selective pressure during strain construction and maintenance. Furthermore, these strains provide defined points of minicell induction that improve minicell purification while creating conditions that allow strain manipulation prior to, during, and following minicell production. By way of non-limiting example these manipulations may be protein production that the cytoplasmic redox state, modify plasmid copy number, and/or produce chaperone proteins.

[1145] For minicell production, a minicell producing strain described in the previous section is grown overnight in Luria broth (LB) supplemented with 0.1% dextrose, 100  $\mu\text{g/ml}$  ampicillin, and when using the single-copy *ftsZ* construct, 15  $\mu\text{M}$  IPTG. All incubations were performed at 37° C. For minicell induction only, overnight strains are subcultured  $\frac{1}{1000}$  into the same media. If minicell induction is to be coupled with co-expression of other proteins that are controlled by a catabolite repression-sensitive regulator, dextrose was excluded. Minicell induction is sensitive to aeration and mechanical forces. Therefore, flask size, media volume and shake speed is critical for optimal yields. Likewise, bioreactor conditions must be properly regulated to optimize these production conditions.

[1146] In shake-flask cultures, strains are grown to early exponential (log) phase as monitored by optical density (OD) at 600 nm ( $\text{OD}_{600}$  0.05-0.20). (Bioreactor conditions may differ significantly depending on the application and

yield desired). For minicell induction alone, early log phase cultures are induced with the appropriate inducer concentration shown in Table 6. For coupled co-expression, these cultures are induced as shown in Table 6 for the appropriate minicell regulator, while the coupled protein(s) is induced with the inducer appropriate for the regulator controlling the synthesis of that protein. Cultures are grown under the appropriate conditions and harvested during late log ( $OD_{600}$  0.8-1.2). Depending on the application, minicell induced cultures may be immediately chilled on ice prior to purification, or maintained at room temperature during the harvesting process.

[1147] To separate minicells from viable, parental cells, cultures are subjected to differential centrifugation (Voros, J., and R. N. Goodman. 1965. Filamentous forms of *Erwinia amylovora*. *Phytopathol.* 55:876-879). Briefly, cultures are centrifuged at 4,500 rpm in a GSA rotor for 5 min. Supernatants are removed to a fresh bottle and centrifuged at 8,000 rpm for an additional 10 min to pellet minicells. Pelleted minicells (containing contaminating parental cells) are resuspended in 2 ml LB, LBD (LB supplemented with 0.1% dextrose), Min (minimal M63 salt media) (Roozen, K. J., et al. 1971. Synthesis of ribonucleic acid and protein in plasmid-containing minicells of *Escherichia coli* K-12. *J. Bacteriol.* 107:21-23), supplemented with 0.5% casamino acids) or MDT (minimal M63 salt media, supplemented with 0.5% casamino acids, 0.1% dextrose, and thiamine). Resuspended minicells are next separated using linear density gradients. By way of non-limiting example, these gradients may contain sucrose (Cohen A., et al. 1968. The properties of DNA transferred to minicells during conjugation. *Cold Spring Harb. Symp. Quant. Biol.* 33:635-641), ficol, or glycerol. For example, linear sucrose gradients range from 5-20% and are poured in LB, LBD, Minor MDT. Using a SW28 swinging bucket rotor, gradients are centrifuged at 4,500 rpm for 14 min. Banded minicells are removed, mixed with LB, LBD, Minor MDT, and using a JA-20 rotor are centrifuged at 13,000 rpm for 12 min. Following centrifugation, pellets are resuspended in 2 ml LB, LBD, Minor MDT and subjected to a second density gradient. Following the second density separation, banded minicells are removed from the gradient, pelleted as described, and resuspended in LB, LBD, Minor MDT for use and/or storage.

[1148] Purified minicells are quantitated using an  $OD_{600}$  measurement as compared to a standard curve incorporating LPS quantity, minicell size, and minicell volume. Quantitated minicells mixtures are analyzed for contaminating, viable parental cells by plating on the appropriate growth media (Table 9).

TABLE 9

MINICELL PURIFICATION AND PARENTAL CELL QUANTITATION				
Purification	Total cells	Total parental cells	MC/PC ratio	Fold-purification
Before	$4.76 \times 10^{11}$	$3.14 \times 10^{11}$	0.25/1	—
After	$1.49 \times 10^{11}$	$6.01 \times 10^{11}$	$2.48 \times 10^6/1$	$5.23 \times 10^6$

## Example 14

## Protoplast Formation

[1149] In order to allow a membrane receptor to be presented to the outside environment (displayed), minicells

are made into protoplasts. In order to make the integral membrane protein receptors in the inner membrane more accessible for ligand binding, the outer membrane and cell wall were removed. The removal of the outer membrane and cell wall from *E. coli* whole cells and minicells to produce protoplasts was performed essentially according to previously described protocols with a few modifications (Birdsell et al., Production and Ultrastructure of Lysozyme and Ethylenediaminetetraacetate-Lysozyme Spheroplasts of *Escherichia coli*, *J. Bacteriol.* 93:427-437, 1967; Weiss et al., Protoplast Formation in *Escherichia Coli*, *J. Bacteriol.* 128:668-670, 1976. Both minicells and whole cells were processed the same way.

[1150] In brief, the cells were grown to mid-log phase and pelleted at room temperature (minicells were isolated from cultures in mid-log phase). The pellet was washed twice with 10 mM Tris. Following the second wash protoplast production may be performed using two approaches. In the first approach, following the second wash, the cells were resuspended in 100 mM Tris (pH 8.0) that contained 6-20% sucrose and put in a 37° C. waterbath (the Tris/sucrose buffer was pre-warmed to 37° C.). The volume used to resuspend the cells was determined by the following equation: (volume of cells  $\times OD_{450}$ )/10=resuspension volume. After a 1 minute incubation, 2 mg/mL lysozyme was added to a final concentration of 5-100  $\mu\text{g}/\text{mL}$ . The samples were then incubated for 12 minutes at 37° C. while being gently mixed. Next, 100 mM EDTA (pH 7) was slowly added over a period of 2.5 minutes (amount of EDTA added= $\frac{1}{60}$ - $\frac{1}{10}$  volume of cells) followed by a 10 min incubation at 37° C. The protoplasts are also diluted from 20% sucrose down to either 10% or 5% sucrose, which facilitates the complete removal of the outer membrane and cell wall. The protoplasts thus generated were separated from the outer membrane and cell wall using a sucrose step gradient. A sucrose step gradient does not have a gradual increase in sucrose percentage; rather, it goes directly from one percent to the other. For example, protoplasts generated from whole cells are loaded on a step gradient that is made from 5% and 15% sucrose. The protoplasts spin through the 15% sucrose but the debris generated when making the protoplasts does not spin through the 15% sucrose. The protoplasts are thus separated from the debris. The second method to prepare protoplasts, following the second wash,  $1 \times 10^9$  cells were resuspended with 50 mM Tris, pH 8.0 containing 0.5-50 mM EDTA and 6-20% sucrose. This mixture was incubated at 37° C. for 10 min. Following incubation, the mixture was centrifuged at 13,200 RPM in a microcentrifuge for 2 min. After centrifugation, the pellet was resuspended in 50 mM Tris, pH 8.0 containing 5-100  $\mu\text{g}/\text{ml}$  lysozyme and 6-20% sucrose. This mixture was incubated at 37° C. for 10 min. Following incubation, the mixture was centrifuged at 13,200 RPM in a microcentrifuge for 2 min, resuspended in 50 mM Tris pH 8.0 containing 6-20% sucrose for use.

[1151] An alternative method to remove contaminating LPS is to use affinity absorption with an anti-LPS antibody (Cortex). To accomplish this, the anti-LPS antibody was coated on either an activated agarose or sepharose matrix (Sigma) or epoxy-coated magnetic M-450 beads (Dyna). The spheroplast/protoplast mixture was subjected to the antibody coated matrix either in batch or using column chromatographic techniques to remove contaminating LPS. Following exposure, the unbound fraction(s) was collected and re-exposed to fresh matrix. To monitor the efficiency of

the protoplasting reaction and LPS removal, three constructs were used (Table 10).

TABLE 10

PROTOPLAST MONITORING CONSTRUCTS					
Construct	SEQ		SEQ		Inducer
	ID NO	Plasmid	ID NO	Inducible protein	
PMPX-5	6	pMPX-32	7	$\Delta$ phoA	Rhamnose
PMPX-5	6	pMPX-53	8	phoA	Rhamnose
PMPX-5	6	pMPX-33	9	toxR-phoA	Rhamnose

[1152]

TABLE 11

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 10 CONSTRUCTS		
SEQ ID NO.:	Primer name	5' to 3' sequence
59	AphoA-1	GCCTGTTCTGGAAAACCGGGCTGCTCAGGG
60	AphoA-2	GCGGCTTTCATGGTGTAGAAGAGATCGG
61	AphoA-1-PstI	CCGCGCTGCAGATGCCTGTTCTGGAAAACCGGGCTGCTCAGGG
62	AphoA-2-XbaI	GCGCCTCTAGATTATTATTTTCAGCCCCAGAGCGGCTTTCATG GTGTAGAAGAGATCGG
63	PhoA-1	GTCACGGCCGAGACTTATAGTCGC
64	PhoA-2	GCGGCTTTCATGGTGTAGAAGAGATCGG
65	PhoA-1-PstI	CCGCGCTGCAGATGTCACGGCCGAGACTTATAGTCGC
66	PhoA-2-XbaI	GCGCCTCTAGATTATTATTTTCAGCCCCAGAGCGGCTTTCATG GTGTAGAAGAGATCGG
67	T-phoA-1-PstI	CCGCGCTGCAGATGAACTGGGGAATCGACTGTTTATTCTGA TAGCGGTCTTACTTCCCTCGCAGTATTACTGCTCATGCCTG TTCTGGAAAACCGGGCTGCTCAGGG
68	T-phoA-2-XbaI	GCGCCTCTAGATTATTATTTTCAGCCCCAGAGCGGCTTTCATG GTGTAGAAGAGATCGG

[1153] Oligonucleotides SEQ ID NOS.:59, 60, 61 and 62 were used to amplify phoA lacking a leader sequence ( $\Delta$ phoA) from the *E. coli* chromosome. Once amplified, this region was inserted into SEQ ID NO.: 6 using PstI and XbaI to create SEQ ID NO.: 7.

[1154] Oligonucleotides SEQ ID NOS.:63, 64, 65 and 66 were used to amplify phoA containing a leader sequence (phoA) from the *E. coli* chromosome. Once amplified, this region was inserted into SEQ ID NO.: 6 using PstI and XbaI to create SEQ ID NO.: 8.

[1155] Oligonucleotides SEQ ID NOS.:59, 60, 67 and 68 were used to amplify phoA lacking a leader sequence ( $\Delta$ phoA) from the *E. coli* chromosome and form a translational fusion between the transmembrane domain of toxR from *Vibrio cholerae*. Once amplified, this region was inserted into SEQ ID NO.: 6 using PstI and XbaI to create SEQ ID NO.: 9.

[1156] By co-expression of minicells and protein, minicells were prepared that contained cytoplasmic PhoA (pMPX-32 expresses phoA lacking a leader sequence [ $\Delta$ phoA]), periplasmic PhoA (pMPX-53 expresses native phoA that exports to the periplasmic space), or inner mem-

brane-bound PhoA (pMPX-33 expresses phoA lacking a leader sequence fused to the transmembrane domain (TMD) of the toxR gene product from *Vibrio cholerae*). Using these expressed proteins, the efficiency of minicell protoplasting was monitored (Table 12).

TABLE 12

EFFICIENCY OF MINICELL PROTOPLAST PREPARATION AND PURIFICATION					
Step	Location <sup>a</sup>	$\Delta$ PhoA	PhoA	T-PhoA	LPS total <sup>b</sup>
Minicell	Pellet	100	100	100	100
EDTA/lysozyme	Whole	100	100	100	100

TABLE 12-continued

EFFICIENCY OF MINICELL PROTOPLAST PREPARATION AND PURIFICATION					
Step	Location <sup>a</sup>	$\Delta$ PhoA	PhoA	T-PhoA	LPS total <sup>b</sup>
1 <sup>st</sup> Anti-LPS	Pellet	80	0	80	30
2 <sup>nd</sup> Anti-LPS	Pellet	60	0	60	0

<sup>a</sup> Measuring the location of protein being measured using an anti-BAP antibody (Sigma). Pellet refers to the presence of the expressed protein in the low-speed centrifugation pellet. These pellets contain only intact cellular bodies. Whole refers to the reaction mixture prior to low-speed centrifugation.

<sup>b</sup> Measured using a slot-blot apparatus (Bio-Rad) using the anti-LPS antibody (Cortex)

[1157] The data suggests that periplasmic PhoA is lost during the preparation, while both cytoplasmic and membrane-bound PhoA are retained in a cellular body that lacks LPS. However, during this process ~40% of the total minicell content is lost.

## Example 15

## T7-Dependent Induction of Expression

[1158] Expression from the pCAL-c expression vector is driven from a T7 bacteriophage promoter that is repressed by the LacI gene product. Transcription of the DNA into mRNA, and subsequent translation of mRNA into proteins, does not occur as long as the LacI repressor is bound to the T7 promoter. However, in the presence of IPTG, the LacI repressor does not bind the T7 promoter. Thus, induction of expression from pCAL-c sequences is dependent on the presence of IPTG. Slightly different protocols were used for the induction of *Escherichia coli* whole and for the induction of minicells. Slight differences are also present in the protocols for induction of minicells for <sup>35</sup>S-methionine labeling of proteins in contrast to those for the induction of minicells for Western blot analysis. These induction protocols are described below.

[1159] For expression in *E. coli* whole cells, the cells were first grown overnight in 3 mL of LB and antibiotics. The cultures were screened for the presence of the desired expression element as previously described. Cultures containing the desired expression elements were diluted 1:100 and grown to an OD<sub>600</sub> of between 0.4 to 0.6. The culture size varied depending on the intended use of the cells. IPTG was then added to a final concentration of 200 μg/mL, and the cells were shaken at 30° C. for 4 hours. Following the induction, cells were harvested for analysis.

[1160] The induction of minicells was carried out as follows. The minicells were diluted in MMM buffer to 1 mL total volume according to the concentration obtained from the isolation procedure (OD<sub>450</sub> of about 0.5). The cells were then treated with 50 μg/mL of cycloserine for 30 minutes at 37° C. to stop whole cell growth. Following the cycloserine treatment the cells were provided with an amino acid, methionine, which the MMM buffer does not contain. For <sup>35</sup>S-labeled protein induction <sup>35</sup>S-methionine was added to the minicell sample whereas, for unlabeled protein induction unlabeled methionine was added. Fifteen (15) μCi of <sup>35</sup>S-methionine (Amersham Pharmacia Biotech, Piscataway, N.J.) was added to the samples for radiolabeling and 5 μmol of methionine was added to the non-labeled minicell samples. Two hundred (200) μg/mL IPTG was also added to the minicell samples, which were then shaken at 30° C. for about 4 hours. Following induction, the minicells were harvested for further preparation or analysis.

## Example 16

## Western Blot Analysis

[1161] The CBP detection kit was purchased from Stratagene. SDS running buffer, 10% Tris-HCl ready gels, Kaleidoscope Pre-stained Standards, and Laemmli Sample Buffer were purchased from BIO RAD (Hercules, Calif.). GFP (FL) HRP antibodies were purchased from Santa Cruz Biotechnology (Santa Cruz, Calif.). Edg-3CT antibody an antibody directed to the carboxy terminus of was purchased from Exalpha Biologicals (Boston, Mass.). Anti-6× His antibody, positrope, and the WesternBreeze Kit were purchased from Invitrogen (Carlsbad, Calif.). Protocols were carried out essentially according to the manufacturer's instructions unless otherwise indicated.

[1162] Three different Western blot protocols were used to detect protein expression in both a minicell expression system and in a whole cell expression system. For both systems, the SDS-PAGE gel and the transfer protocols were essentially as follows. The samples were denatured by diluting the samples 1:1 in Laemmli buffer (BIORAD) and then sonicated for 10 mm. The denatured samples were loaded onto a 10% Tris-Glycine gel (BIORAD) and electrophoresed at 130 V for about 1.5 hours in 1× SDS running buffer (BIORAD). The electrophoresed proteins were electrotransferred to nitrocellulose membranes at 0.5 Amps for 1.5 hours in Transfer Buffer (5.8 g Tris, 2.9 g glycine, 200 mL methanol, and 3.7 mL of 10% SDS). The nitrocellulose membranes comprising the transferred proteins were used for Western blotting.

[1163] GFP Western blots were carried out as follows. The nitrocellulose membrane was blocked for 2 hours with 5% milk in PBST (PBS buffer with 0.05% Tween). Following the blocking step the nitrocellulose membrane was washed twice with PBST. For the detection of GFP protein, an anti-GFP-HRP conjugated antibody (Santa Cruz Biotechnology) was used at a dilution of 1:3000 in PBST (HRP, horse radish peroxidase). The nitrocellulose membrane was incubated in the anti-GFP-HRP antibody solution for one hour and then washed twice with PBST. GFP proteins on the nitrocellulose membrane were detected and visualized using the ECL system (Amersham).

[1164] The His-tagged Edg-1 and Edg-3 proteins were detected using a mouse anti-6× His antibody from Invitrogen and the WesternBreeze chemoluminescent Kit (Invitrogen). The antibody was diluted 1:4000 in buffers provided by the WesternBreeze Kit. The WesternBreeze immunoblot was carried out essentially according to the manufacturer's protocol. The Edg-1-CBP and GFP-CBP fusion proteins were detected using the CBP detection Kit (Stratagene). All antibodies and substrates were provided in the Kit. **FIG. 3** is a photo of the Western hybridization results showing the presence of Edg-1-6× His and Edg-3-6× His in minicells and parent cells.

## Example 17

## Methods to Induce Expression

[1165] Expression in minicells may proceed following purification of minicells and/or minicell protoplasts from parental cells and LPS constituents, respectively. However, for some applications it is suitable to co-express proteins of interest with minicell induction. For these approaches, one may use the protocol described in EXAMPLE 13 for expression of the phoA constructs. By way of non-limiting example, either of these approaches may be accomplished using one or more of the following expression constructs (Table 13).

TABLE 13

EXPRESSION CONSTRUCTS				
Plasmid	Regulatory element(s)	inducer	Plasmid	SEQ ID NO.:
pMPX-5	rhaRS	Rhamnose	pUC-18	6
pMPX-7	uidR	β-glucuronate	pUC-18	10
pMPX-8	melR	Melibiose	pUC-18	11

TABLE 13-continued

EXPRESSION CONSTRUCTS				
Plasmid	Regulatory element(s)	inducer	Plasmid	SEQ ID NO.:
pMPX-18	araC	Arabinose	pUC-18	12
pMPX-6	araC	Arabinose	pUC-18	13

[1166]

region from the *E. coli* chromosome. Once amplified, this region was inserted into pUC 18 using HindIII and SalI to create SEQ ID NO.: 11.

[1170] Oligonucleotides SEQ ID NOS.:81, 82, 83 and 84 were used to amplify the araC gene and its divergent control region from the *E. coli* chromosome. Once amplified, this region was inserted into pUC18 using HindIII and PstI to create SEQ ID NO.: 12.

[1171] Oligonucleotides SEQ ID NOS.:81, 82, 85 and 86 were used to amplify the araC gene and its divergent control

TABLE 14

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 13 CONSTRUCTS		
SEQ ID NO.:	Primer name	5' to 3' sequence
69	Rha-1	GCGAATTGAGATGACGCCACTGGC
70	Rha-2	CCTGCTGAATTCATTAAACGACCAG
71	Rha-1-HindIII	CGCGCAAGCTTAATTAATCTTTCTGCGAATTGAGATGACGCCACTGGC
72	Rha-2-PstI	CGCCGTAATCGCCGCTGCAGAATGTGATCCTGCTGAATTCATTAACGACCAG
73	Uid-1	CGCAGCGCTGTTCCCTTTGCTCG
74	Uid-2	CCTCATTAAAGATAATAATACTGG
75	Uid-1-HindIII	GCCGCAAGCTTCGCAGCGCTGTTCCCTTTGCTCG
76	Uid-2-PstI	CCAATGCATTGGTTCTGCAGGACTCCTCATTAAAGATAATAATACTGG
77	Mel-1	CGTCTTTAGCCGGGAAACG
78	Mel-2	GCAGATCTCCTGGCTTGC
79	Mel-1-HindIII	GCCGCAAGCTTCGTCTTTAGCCGGGAAACG
80	Mel-2-SalI	CGGTGACGCAGATCTCCTGGCTTGC
81	Ara-1	CAAGCCGTCAATTGTCTGATTTCG
82	Ara-2	GGTGAATTCCTCTGCTAGCCC
83	Ara-1-HindIII	GCGCCAAGCTTCAAGCCGTCAATTGTCTGATTTCG
84	Ara-2-PstI	CTGCAGGGTGAATTCCTCTGCTAGCCC
85	Ara-1-XhoI	GCTTAACTCGAGCTTAATAACAAGCCGTCAATTGTCTGATTTC
86	Ara-2-SstI	GCTTAAACCGGGCCAAGCTTGCATGCCTGCTCC

[1167] Oligonucleotides SEQ ID NOS.:69, 70, 71 and 72 were used to amplify the rhaRS genes and their divergent control region from the *E. coli* chromosome. Once amplified, this region was inserted into pUC18 using HindIII and PstI to create SEQ ID NO.: 6.

[1168] Oligonucleotides SEQ ID NOS.:73, 74, 75 and 76 were used to amplify the uidR control region, the uidR gene and the control region for expression from the *E. coli* chromosome. Once amplified, this region was inserted into pUC18 using HindIII and PstI to create SEQ ID NO.: 10.

[1169] Oligonucleotides SEQ ID NOS.:77, 78, 79 and 80 were used to amplify the meiR gene and its divergent control

region was PCR amplified from pBAD-24. Once amplified, this region was inserted into pEGFP (Clontech) using XhoI and SstI to create SEQ ID NO.: 13.

[1172] Except of pMPX-6, these expression constructs contain the same multiple cloning site. Therefore, any protein of interested may be inserted in each modular expression construct for simple expression screening and optimization.

[1173] By way of non-limiting example, other proteins that may be expressed are listed in Table 15.



TABLE 15

OTHER EXPRESSED PROTEINS				
Protein	Origin	Construct	Purpose	SEQ ID NO.:
Edg3	Rat	native	GPCR	14
$\beta$ 2AR	Human	native	GPCR	15
TNFR-1a (human)	Human	residues 29-455	Receptor	18
TNFR-1b (human)	Human	residues 41-455	Receptor	17
TNF (human)	Human	native	Gene transfer	19

TABLE 15-continued

OTHER EXPRESSED PROTEINS				
Protein	Origin	Construct	Purpose	SEQ ID NO.:
T-EGF	Human	chimera	Gene transfer	20
T-Invasin	<i>Y. pseudotuberculosis</i>	chimera	Gene transfer	21

[1174]

TABLE 16

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 15		
SEQ ID NO.:	Primer name	5' to 3' sequence
87	Edg-1	GGCAACCACGCACGCGCAGGGCCACC
88	Edg-2	CAATGGTGATGGTGATGATGACCGG
89	Edg-1-SalI	CGCGGTCGACATGGCAACCACGCACGCGCAGGGCCACC
90	Edg-2-KpnI	GCGCCGGTACCTTATCAATGGTGATGGTGATGATGACCGG
91	$\beta$ 2AR-1	GGGGCAACCCGGGAACGGCAGCGCC
92	$\beta$ 2AR-2	GCAGTGAGTCATTTGTACTACAATTCCCTCC
93	$\beta$ 2AR-1-SaII	CGCGGTCGACATGGGGCAACCCGGGAACGGCAGCGCC
94	$\beta$ 2AR-2-BamHI	GCGCCGGATCCTTATTATAGCAGTGAGTCATTTGTACTACAATTCCCTCC
95	TNFR(29)-1	GGACTGGTCCCTCACCTAGGGGACAGGG
96	TNFR(29)-2	CTGAGAAGACTGGGCGGGGGGGAGG
97	TNFR(29)-1-SalI	CGCGGTCGACATGGGACTGGTCCCTCACCTAGGGGACAGGG
98	TNFR(29)-2-KpnI	GCGCCGGTACCTTATTACTGAGAAGACTGGGCGGGGGGGAGG
99	TNFR(41)-1	GATAGTGTGTGTCCCC
100	TNFR(41)-2	CTGAGAAGACTGGGCGC
101	TNFR(41)-1-NcoI	GGGAGACCATGGATAGTGTGTGTGTCCCC
102	TNFR(41)-2-XbaI	GCCTCATCTAGATTTACTGAGAAGACTGGGCGC
103	TNF-1	GAGCACTGAAAGCATGATCCGGGACG
104	TNF-2	CAGGGCAATGATCCCAAAGTAGACCTGC
105	TNF-1-EcoRI	CCGCGGAATTCATGAGCACTGAAAGCATGATCCGGGACG
106	TNF-2-HindIII	GGCGCAAGCTTATCACAGGGCAATGATCCCAAAGTAGACCTGC
107	T-EGF-1	TCTGATAGCGGTCTTACTTCCCCTCGCAGTATTACTGCTCAATAGTGACTCTGAATGTCCCCTGTCCCACGATGGGTACTGCCTCCATGATGGTGTGTGCATGTATATTG
108	T-EGF-2	AGGTCTCGGTACTGACATCGTCCCCTGATGTAGCCAACAACACAGTTGCATGCATACTTGTCCAATGCTTCAATATACATGCACACACCATCATGGAGGCA

TABLE 16-continued

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 15		
SEQ ID NO.: Primer name	5' to 3' sequence	
109	T-EGF-3	CCGCGGGTACCATGAACTTGGGAATCGACTGTTTATTCT GATAGCGGTCTTACTTCCCTCG
110	T-EGF-4	GCGCCAAGCTTATTAGCGAGTCCACCACCTTCAGGTCT CGGTACTGACATCGCTCCCG
111	Inv-1	TCATTCACATTGAGCGTCACCG
112	Inv-2	TTATATTGACAGCGCACAGAGCGG
113	Inv-1-ToxR-EcoRI	GCAAGAATTTCCATGAACTTGGGAATCGACTGTTTATT CTGATAGCGGTCTTACTTCCCTCGCAGTATTACTGCTCT CATTACATTGAGCGTCACCG
114	Inv-2-PstI	CGCGGTTACGTAAGCAACTGCAGTTATATTGACAGCGCA CAGAGCGG

[1175] Oligonucleotides SEQ ID NOS.:87, 88, 89 and 90 were used to amplify rat Edg3 from rat cDNA. Once amplified, this region was inserted into SEQ ID NO.: 6 (pMPX-5) using Sall and KpnI to create SEQ ID NO.: 14.

[1176] Oligonucleotides SEQ ID NOS.:91, 92, 93 and 94 were used to amplify human  $\beta$ 2 adrenergic receptor ( $\beta$ 2AR) from human heart cDNA. Once amplified, this region was inserted into SEQ ID NO.: 6 (pMPX-5) using Sall and BamHI to create SEQ ID NO.: 15.

[1177] Oligonucleotides SEQ ID NOS.:95, 96, 97 and 98 were used to amplify human tumor necrosis factor receptor (TNFR residues 29-455) from human Jurkat CL71 cDNA. Once amplified, this region was inserted into SEQ ID NO.: 12 (pMPX-18) using Sall and KpnI to create SEQ ID NO.:18.

[1178] Oligonucleotides SEQ ID NOS.:99, 100, 101 and 102 were used to amplify human tumor necrosis factor receptor (TNFR residues 41-455) from human Jurkat CL71 cDNA. Once amplified, this region was inserted into pBAD24 using NcoI and XbaI to create SEQ ID NO.: 17.

[1179] Oligonucleotides SEQ ID NOS.:103, 104, 105 and 106 were used to amplify human tumor necrosis factor (TNF) from human Jurkat CL71 cDNA. Once amplified, this region was inserted into SEQ ID NO.: 13 (pMPX-6) using EcoRI and HindIII to create SEQ ID NO.:19.

TABLE 17

PROGRAM TO ANNEAL GRADIENT PCR WITH PFX POLYMERASE		
Step	Temp (° C.)	Time (min)
1	95	2.0
2	95	0.5
3	64	0.5
4	68	2.5
5	Goto 2, 2X	
6	95	0.5
7	62	0.5
8	68	2.5
9	Goto 6, 4X	

TABLE 17-continued

PROGRAM TO ANNEAL GRADIENT PCR WITH PFX POLYMERASE		
Step	Temp (° C.)	Time (min)
10	95	0.5
11	60	0.5
12	68	2.5
13	Goto 10, 6X	
14	95	0.5
15	58	0.5
16	68	2.5
17	Goto 14, 24X	
18	4	hold
19	end	

[1180] Oligonucleotides SEQ ID NOS.: 107, 108, 109 and 110 were mixed and PCR amplified using anneal gradient PCR (Table 17) to form mature human epidermal growth factor (EGF) (residues 971-1023) translationally fused to the transmembrane domain of toxR from *Vibrio cholerae*. Once amplified, this region was inserted into SEQ ID NO.: 13 (pMPX-6) using KpnI and HindIII to create SEQ ID NO.:20.

[1181] Using PFX polymerase (Invitrogen) oligonucleotide SEQ ID NOS.:111, 112, 113 and 114 were used to amplify invasin residues 490-986 (inv) from *Yersinia pseudotuberculosis* chromosomal DNA and form a translational fusion between the transmembrane domain of toxR from *Vibrio cholerae*. Once amplified, this region was inserted into SEQ ID NO.:13 (pMPX-6) using EcoRI and PstI to create SEQ ID NO.:21.

[1182] These proteins were proof-of-principle constructs used to evaluate the minicell platform. For purposes of this initial evaluation, all proteins except TNF, T-EGF and T-Invasin were cloned into pMPX-5, with these later proteins cloned into pMPX-6 for gene transfer experiments.

[1183] Whether the approach for protein expression is co-expression with minicell induction or expression following minicell and/or protoplast isolation, the procedure to transform the expression constructs is the same. To accom-

plish this, protein constructs were initially cloned into *E. coli* MG1655 and then into the minicell producing strain of interest. Transformation events were selected prior to minicell induction. For co-induction of protein and minicells, see the protocol for phoA expression above. For post-minicell and/or protoplast purification induction experiments, following minicell purification and/or protoplast preparation and purification, these cellular bodies were induced for protein production in either LBD or MDT at a minicell or protoplast/volume ratio of  $1 \times 10^9$  minicells or protoplasts/1 ml media. Media was supplemented with the appropriate inducer concentration (see Table 6). Protein induction is sensitive to a variety of factors including, but not limited to aeration and temperature, thus reaction volume to surface area ratio is important, as is the method of shaking and temperature of induction. Therefore, each protein must be treated as required to optimize expression. In addition to expression parameters, protoplasted minicells are sensitive to osmotic and mechanical forces. Therefore, protoplast protein induction reactions must also contain 10% sucrose with greater volume to surface area ratios than required for intact minicells to achieve similar aeration at lower revolutions.

[1184] Using the T-PhoA as a non-limiting example, protein expression was performed during and following minicell isolation. To accomplish this task, t-phoA co-expressed with minicell induction was compared to t-phoA expressed after minicell isolation. In both cases, overnight minicell-producing parental strains containing pMPX-5::t-phoA were subcultured into LBD supplemented with the appropriate antibiotic. Cultures were grown to OD<sub>600</sub> 0.1 and induced for minicell production alone or for both minicell and

although the kinetics are slower for the post-minicell purification induction protocol, the end result is equivalent.

TABLE 18

COMPARATIVE EXPRESSION: CO-EXPRESSION VERSUS POST MINICELL PURIFICATION INDUCTION		
Time of induction	Purified minicell induction <sup>a</sup>	Co-expression induction <sup>a</sup>
1.0	8.0	—
2.0	—	812.2
4.0	70.0	—
14.0	445.0	—

<sup>a</sup>Nanogram expressed T-PhoA per  $1 \times 10^9$  minicells.

[1185] Using the co-expression induction procedure, the amount of membrane-associated T-PhoA was measured and compared for both parental cells and minicells. Briefly, following co-expression induction of T-PhoA and minicells, minicells were purified and their membranes isolated. For membrane isolation, minicells containing expressed T-PhoA were subjected to three rounds of freeze-thaw lysis in the presence of 10  $\mu$ g/ml lysozyme. Following freeze-thaw cycling, the reaction was subjected to sonication. Sonicated material was centrifuged at 6,000 rpm in a microcentrifuge for 5 min at room temperature. Supernatants were transferred to a fresh 1.5 ml Eppendorf tube and centrifuged at 70,000 rpm using a TLA-100 rotor. Following centrifugation, the pellet was resuspended in buffer and analyzed for total T-PhoA protein (Table 19) and T-PhoA enzyme activity (Table 20).

TABLE 19

MEMBRANE ASSOCIATED T-PHOA: PARENTAL CELLS VERSUS MINICELLS						
Cell type <sup>a</sup>	Protein total <sup>a</sup>	T-PhoA total <sup>b</sup>	T-PhoA % total	Protein membrane associated <sup>a</sup>	T-PhoA membrane associated <sup>b</sup>	T-PhoA % membrane protein total
Parental cells	107.5	5.3	4.9	10.7	3.1	29.0
Minicells	4.6	0.8	17.5	1.0	0.5	50.0
Minicells EQ <sup>b</sup>	25.2	4.4	—	5.5	2.7	—

<sup>a</sup> Total protein as determined by BCA assay (Pierce)

<sup>b</sup> Microgram expressed T-PhoA per  $1 \times 10^9$  minicells as determined via Western using an anti-PhoA antibody (Sigma) versus a PhoA standard curve (BCA determined).

<sup>c</sup> Equivalent membrane lipid to parental cell

protein production. Both cultures were harvested at OD<sub>600</sub> 1.0 and minicells produced were harvested as described above. Minicells to be induced for T-phoA production following purification were induced by introducing  $1 \times 10^9$  purified minicells into a 15 ml culture tube containing 1 ml MDT with 1 mM L-rhamnose. Minicell protein induction was allowed to proceed for up to 14 hours and compared to protein production obtained using the co-expression approach. For each approach, minicells were fractionated and analyzed for membrane association, total protein, and membrane association-dependent enzymatic activity. These observations were compared to post-induction, pre-isolation parental cell/minicell (PC/MC) mixtures from the co-expressed reactions. The first observation was that co-expression of minicell and protein induction was superior to post-minicell purification induction (Table 18). However,

[1186]

TABLE 20

PHOA ENZYMATIC ACTIVITY <sup>a</sup> (RELATIVE UNITS): PARENTAL CELLS VERSUS MINICELLS.			
Cell type <sup>b</sup>	Unlysed	Lysed, total	Lysed, membrane
Parent cell	—	358	240
Minicell	275	265	211
Minicell EQ <sup>c</sup>	1,504	1,447	1,154

<sup>a</sup>Activity determined colorimetrically using PNPP measuring optical density at 405 nm

<sup>b</sup>Based on  $1 \times 10^9$  parental cells or minicells per reaction

<sup>c</sup>Equivalent membrane lipid to parental cell

[1187] These results suggest that co-expression induction of T-PhoA and minicells together results in minicells containing an equivalent amount of T-PhoA produced in both parental cells and minicells. However, the percent of T-PhoA compared to total protein is 3.5× greater in minicells than in parental cells. Furthermore, of the protein made, T-PhoA constitutes 50% of the total membrane protein in minicells, whereas it is only 29% in parental cells. It should be noted that the T-PhoA protein associated with the membrane can be easily removed by treatment with mild, non-ionic detergent suggesting that the T-PhoA present in the membrane pellet is indeed associated with the membrane and not an insoluble, co-sedimenting precipitate (data not shown). Finally, PhoA is a periplasmic enzyme that requires export to the periplasmic space for proper folding and disulfide bond formation. Both of which are required for enzymatic activity. In the time course of this experiment, expression of APhoA lacking a leader sequence does not demonstrate enzymatic activity. Furthermore, there is no difference between unlysed and lysed minicells containing expressed T-PhoA (Table 20) also demonstrating that the PhoA enzyme domain of the T-PhoA chimera must be present in the periplasmic space. Therefore, the T-PhoA construct must membrane associate and the PhoA domain must orient into the periplasmic space for enzymatic activity. Thus, when comparing equivalent amounts of membrane lipid between parental cells and minicells in Table 20, membrane association-dependent T-PhoA activity is almost 5× greater than in parental cells. Taking into account the data in Table 19

chromosome, it is also possible that the overexpression of this protein is readily finding membrane-binding sites in the absence of chromosomally produced competitors present in parental cells. Furthermore, overexpression of proteins often leads to increased protease expression. Because minicells do not contain chromosome, these otherwise degraded surplus T-PhoA is allowed the continued opportunity to insert and properly fold in the membrane, an attribute that could lead favor to overexpression of more complex membrane proteins.

#### Example 18

##### Exemplary Methods to Induce and Study Complex Membrane Proteins

[1188] Expression of non-native (exogenous) complex membrane proteins in bacterial systems can be difficult. Using the minicell system, we are able to eliminate toxicity issues. However, issues still remain with proper translation, compartmentalization at the membrane, insertion in the membrane and proper folding for native activity. To account for these potential problems we have constructed a modular chimeric system that incorporates leader sequences and chaperone-recognized soluble domains that are native to our bacterial minicell system. In addition, we created modular constructs that overexpress the native chaperones groESL and trigger factor (tig). Finally, we have constructed minicell-producing strains that contain mutations that effect protein export and disulfide bond formation. For non-limiting examples of these constructs see Table 21.

TABLE 21

NON-LIMITING TOOLS FOR EXOGENOUS COMPLEX PROTEIN SYNTHESIS AND FUNCTION				
Tool	Reference	Residues of sequence	Purpose	SEQ ID NO
pMPX-5::phoA leader	—	1-48	Membrane targeting	22
pMPX-5::phoA leader	—	1-494	Membrane targeting	23
pMPX-5::malE leader	1	1-28	Membrane targeting	24
pMPX-5::malE leader	1	1-370	Membrane targeting	25
pMPX-17 (groESL, tig)	—	—	Chaperone	26
pMPX-5::trxA::FLAG	2	2-109 <sup>a</sup>	Solubility	27

<sup>a</sup>Residues do not include FLAG sequence.

where 50% of T-PhoA is in the membrane compared to 29% in parental cells, the difference in T-PhoA membrane association is not sufficient to explain the almost 5× increase in minicell activity. These observations suggest that minicells contain a capacity to support more expressed membrane protein than parental cells and that the protein that associates with the membrane is more active. This activity may be simply result from minicells allowing greater efficiency of folding and disulfide bond formation for this particular protein. However, do to the fact that minicells do not contain

#### REFERENCES TO TABLE 21

- [1189] 1. Grisshammer, R., et al. 1993. Expression of a rat neurotensin receptor in *Escherichia coli*. *Biochem. J.* 295:571-576.
- [1190] 2. Tucker, J., and R. Grisshammer. 1996. Purification of a rat neurotensin receptor expressed in *Escherichia coli*. *Biochem. J.* 317:891-899.

TABLE 22

<u>OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 21 CONSTRUCTS</u>		
SEQ ID NO.:Primer name	5' to 3' sequence	
115	PhoA lead-1	GTCACGGCCGAGACTTATAGTCGC
116	PhoA lead-2	GGTGTCCGGGCTTTGTACAGG
117	PhoA lead-1-PstI	CGCGGCTGCAGATGTCACGGCCGAGACTTATAGTCGC
118	PhoA lead-2-XbaI	CGCGGTCTAGATTCTGGTGTCCGGGCTTTGTACAGG
119	PhoA complete	CAGCCCCAGAGCGGCTTTTCATGG
120	PhoA complete-2-XbaI	CGCGGTCTAGATTTTCAGCCCCAGAGCGGCTTTTCATGG
121	MalE lead-1	CGCGGCTGCAGATGAAAATAAAAACAGGTGCACGC ATCCTCGCATTATCCGCATTAACGACGATGATGTTTT CCGCCTCGGCTCTCGCCAAAATCTCTAGACGCGG
122	MalE lead-2	CCGCGTCTAGAGATTTTGGCGAGAGCCGAGGCGGAA AACATCATCGTCGTTAATGCGGATAATGCGAGGATG CGTGCACCTGTTTTATTTTCATCTGCAGCCGCG
123	MalE-1	GGTGCACGCATCCTCGCATTATCCGC
124	MalE-2	CGGCATACCAGAAAGCGGACATCTGC
125	MalE-1-PstI	CGCGGCTGCAGATGAAAATAAAAACAGGTGCACGC ATCCTCGCATTATCCGC
126	MalE-2-XbaI	CGCGGTCTAGAACGCACGGCATAACCAGAAAGCGGACATCTGC
127	Tig-1	CGCGACAGCGCGCAATAACCGTTCTCG
128	Tig-2	GCTGGTTCATCAGCTCGTTGAAAGTGG
129	Tig-1-NarI	GCGCCGGCGCCATACGCGACAGCGCGCAATAACCGTTCTCG
130	Tig-2-XbaI	GGCGCTCTAGATTATTATTACGCTGCTGGTTCATCA GCTCGTTGAAAGTGG
131	Gro-1	GGTAGCACAATCAGATTCGCTTATGACGG
132	Gro-2	GCCGCCATGCCACCCATGCCGCC
133	Gro-1-XbaI	GCGTCTAGAGGTAGCACAATCAGATTCGCTTATGACGG
134	Gro-2-HindIII	GGCGCAAGCTTATTATTACATCATGCCGCCATGCC ACCCATGCCGCC
135	TrxA-1	GCGATAAAATTTATTCACCTGACTGACG
136	TrxA-2	GCGTCGAGGAACCTTTCAACTGACC
137	TrxA-1-Fxa-PstI	CGCGGCTGCAGATGATCGAAGCCCGCTCTAGACTCG AGAGCGATAAAATTTATTCACCTGACTGACG
138	TrxA-2-FLAG-BamHI	CCGCGGGATCCTTATTAATCATCATGATCTTTATAAT CGCCATCATGATCTTTATAATCTCGAGCGCCAGGTT AGCGTCGAGGAACCTTTCAACTGACC

[1191] Oligonucleotides SEQ ID NOS.:115, 116, 117 and 118 were used to amplify the *phoA* leader (residues 1-49) from *E. coli* chromosomal DNA. Once amplified, this region was inserted into SEQ ID NO.: 6 (pMPX-5) using PstI and XbaI to create SEQ ID NO.:22.

[1192] Oligonucleotides SEQ ID NOS.:115, 117, 119 and 120 were used to amplify the complete *phoA* gene from *E. coli* chromosomal DNA. Once amplified, this region was inserted into SEQ ID NO.: 6 (pMPX-5) using PstI and XbaI to create SEQ ID NO.:23.

[1193] Oligonucleotides SEQ ID NOS.:121 and 122 were used to construct the *malE* leader (residues 1-28) sequence. Once annealed, this construct was inserted into SEQ ID NO.: 6 (pMPX-5) using PstI and XbaI to create SEQ ID NO.:24.

[1194] Oligonucleotides SEQ ID NOS.: 123, 124, 125 and 126 were used to amplify the *malE* expanded leader (residues 1-370) from *E. coli* chromosomal DNA. Once amplified, this region was inserted into SEQ ID NO.: 6 (pMPX-5) using PstI and XbaI to create SEQ ID NO.:25.

[1195] Oligonucleotides SEQ ID NOS.:127, 128, 129 and 130 were used to amplify the *tig* control and gene region from *E. coli* chromosomal DNA. Once amplified, this region was ligated to the *groESL* amplified region below using XbaI prior to insertion into SEQ ID NO.: 6 (pMPX-5) using NarI (from the *tig* region) and HindIII (from the *groESL* region) to create SEQ ID NO.:26.

[1196] Oligonucleotides SEQ ID NOS.:131, 132, 133 and 134 were used to amplify the *groESL* control and gene region from *E. coli* chromosomal DNA. Once amplified, this region was ligated to the *tig* amplified region above using XbaI prior to insertion into SEQ ID NO.: 6 (pMPX-5) using NarI (from the *tig* region) and HindIII (from the *groESL* region) to create SEQ ID NO.:26.

[1197] Oligonucleotides SEQ ID NOS.: 135, 136, 137 and 138 were used to amplify *trxA* (residues 2-109) from *E. coli* chromosomal DNA and insert FLAG and Factor Xa sequences. Once amplified, this region was inserted into SEQ ID NO.: 6 (pMPX-5) using PstI and BamHI to create SEQ ID NO.:27.

[1198] By way of non-limiting example, the pMPX-5::phoA leader (residues 1-48), pMPX-5::phoA leader (residues 1-494), pMPX-5::malE leader (residues 1-28), and pMPX-5::malE leader (residues 1-370) constructs are designed to direct expressed exogenous membrane proteins to the minicell cytoplasmic membrane. In addition to these constructs, By way of non-limiting example, mutations in *E. coli* genes *secA* and *secY*, specifically mutation *prlA4* (Strader, J., et al. 1986. Kinetic analysis of lamB mutants suggests the signal sequence plays multiple roles in protein export. *J. Biol. Chem.* 261:15075-15080), permit promiscuous targeting to the membrane. These mutations, like the above constructs are integrated into the minicell expression system. To complement these mutations, the chaperone complex *groESL* and trigger factor have also been incorporated into the expression system. By way of non-limiting example, pMPX-5::trxA::FLAG will be used to create a carboxy-terminal fusion to the protein of interest to increase the membrane insertion efficiency of the membrane protein of interest (Tucker, J., and R. Grishammer. 1996. Purification of a rat neurotensin receptor expressed in *Escherichia*

*coli*. *Biochem. J.* 317:891-899). Also By way of non-limiting example, pMPX-5::FLAG::toxR and pMPX-5::FLAG::XcI constructs will be prepared to create a carboxy-terminal fusion to the protein of interest for use in a reporter-based assay for protein-protein interactions. By way of non-limiting example, the protein of interest for this system is a GPCR. Also By way of non-limiting example, this GPCR may be the neurotensin receptor from rat (Grishammer, R., et al. 1993. Expression of a rat neurotensin receptor in *Escherichia coli*. *Biochem. J.* 295:571-576.), or the  $\beta_2$  adrenergic receptor from humans (Freissmuth, M., et al. 1991. Expression of two  $\beta$ -adrenergic receptors in *Escherichia coli*: functional interaction with two forms of the stimulatory G protein. *Proc. Natl. Acad. Sci.* 88:8548-8552). Insertion of a GPCR into one of these reporter constructs creates a carboxy-terminal fusion between the GPCR of interest and the DNA-binding regulatory domain of the ToxR positive activator, the *kcl* repressor, or the AraC positive activator. To complete this reporter system, By way of non-limiting example pMPX-5::(X)::toxR or pMPX-5::(X)::XcI will be used to create a carboxy-terminal fusion to the protein of interest for use in a reporter-based assay for protein-protein interactions, where (X) may be any protein or molecule involved in an intermolecular or intramolecular interaction. By way of non-limiting example, this molecule of interest may be a G-protein. This G-protein may be the  $G_{\alpha_1}$ -protein from rat (Grishammer, R., and E. Hermans. 2001. Functional coupling with  $G_{\alpha q}$  and  $G_{\alpha i1}$  protein subunits promotes high-affinity agonist binding to the neurotensin receptor NTS-1 expressed in *Escherichia coli*. *FEBS Lett.* 493:101-105), or the  $G_s$ -protein from human (Freissmuth, M., et al. 1991. Expression of two  $\beta$ -adrenergic receptors in *Escherichia coli*: functional interaction with two forms of the stimulatory G protein. *Proc. Natl. Acad. Sci.* 88:8548-8552). Like the GPCR, insertion of a G-protein into one of these reporter constructs creates a carboxy-terminal fusion between the G-protein of interest and the DNA-binding regulatory domain of the ToxR positive activator, the XcI repressor, or other regulatory protein. Finally, these plasmid constructs contain the DNA-binding domain of each regulator; the *ctx* regulatory region from *Vibrio cholerae* (Russ, W. P., and D. M. Engelman. 1999. TOXCAT: a measure of transmembrane helix association in a biological membrane. 96:863-868), or the  $P_{R10R1}$  region of bacteriophage lambda (Hu, J. C., et al. 1990. Sequence requirements for coiled-coils: analysis with lambda repressor-GCN4 leucine zipper fusions. *Science.* 250:1400-1403), respectively. By way of non-limiting example, each binding domain is coupled to a reporter sequence encoding luciferase (Dunlap, P. V., and E. P. Greenberg. 1988. Control of *Vibrio fischeri* lux gene transcription by a cyclic AMP receptor protein-luxR protein regulatory circuit. *J. Bacteriol.* 170:4040-4046), green fluorescent protein (GFP) (Yang, T. T., et al. 1996. Dual color microscopic imagery of cells expressing the green fluorescent protein and a red-shifted variant. *Gene.* 173:19-23; Matthyse, A. G., et al. 1996. Construction of GFP vectors for use in gram-negative bacteria other than *Escherichia coli*. *FEMS Microbiol. Lett.* 145:87-94), or other reporter. Co-expression of these GPCR and G-protein chimeras will create a system measuring the interaction between a GPCR and G-protein within an intact minicell. This system is designed to be used as a positive or negative read-out assay and may be used to detect loss or gain of GPCR function. Although the GPCR-G-protein

interaction is provided as an example, this modular system may be employed with any soluble or membrane protein system measuring protein-protein or other intermolecular interaction.

#### Example 19

##### Exemplary Methods for Gene Transfer Using Minicells or Minicell Protoplasts

[1199] Included in the design of the invention is the use of minicells to transfer genetic information to a recipient cell. By way of non-limiting example, this gene transfer may occur between a minicell and a mammalian cell in vitro, or in vivo, and this gene transfer may occur through cell-specific interactions, through general interactions, or a combination of each. To accomplish this task three basic constructs were created. Each of these constructs is created in pMPX-6 which contains a CMV promoter controlling the synthesis of GFP. The plasmid pMPX-6 was constructed by cloning the *araC* through the multiple cloning site of pBAD24 into pEGFP (Clontech). This construct provided a bacterial regulator as well as a method to monitor the success of gene transfer using GFP expression from the CMV promoter. In design, the protein expressed using the bacterial promoter will drive the cell-cell interaction, while the successful transfer of DNA from the minicell to the recipient cell will initiate the production of GFP. By way of non-limiting example, proteins that will drive the cell-cell interaction may be the invasin protein from *Yersinia pseudotuberculosis*, which stimulates  $\beta 1$  integrin-dependent endocytic events. To properly display the invasin protein on the surface of minicells, the domain of invasin that stimulates these events (residues 490-986) (Dersch, P., and R. R. Isberg, 1999. A region of the *Yersinia pseudotuberculosis* invasin protein enhances integrin-mediated uptake into mammalian cells and promotes self-association. *EMBO J.* 18:1199-1213) was fused to the transmembrane domain of ToxR. Expression of this construct from pMPX-6 will display T-Inv on the surface of the minicell and stimulate endocytosis with any cell displaying a  $\beta 1$  integrin. Thus, T-Inv display will provide a general mechanism of gene transfer from minicells. To provide specificity, By way of non-limiting example, the ligand portion of epidermal growth factor (EGF) may be fused to the transmembrane domain of ToxR, thus creating a protein that will interact with cells displaying the EGF receptor (EGFR). Likewise, tumor necrosis factor (TNF) may also serve this purpose by stimulating cell-cell interactions between minicells displaying TNF and cells displaying TNF receptor (TNFR). Although EGF-EGFR and TNF-TNFR interactions may stimulate cell-cell fusion between minicells and recipient cells, or minicell uptake, this alone may not be sufficient to efficiently transfer genetic information from minicells. Therefore, a genetic approach to increasing the cell-cell genetic transfer may be the development of a genetic switch that senses the specificity interaction, e.g. EGF-EGFR interaction, and turns on the production of a second gene product, e.g. invasin, that stimulates the endocytic event. By way of non-limiting example, this genetic switch may be similar to the GPCR-G-protein interaction reporter system above, in that an extracellular event stimulates the dimerization of a transcriptional active regulator, thus turning on the production of invasin or invasin-like protein. In either approach, the display system to stimulate transfer of genetic information

from minicells to recipient cells may also be applicable to the transfer of substances other than genetic information, e.g. pre-synthesized therapeutic drugs.

[1200] To test this targeting methodology, different pMPX-6 constructs containing each of these general or specific cell-cell interaction proteins will be transformed into a minicell producing strain and either by co-expression induction of minicells, by post-minicell purification induction, or by post-protoplasting induction, minicells displaying the targeting protein of interest will be produced. When using the co-expression induction and post-minicell purification induction of the targeting protein approaches, it is necessary to protoplast the purified minicells after protein induction. Once the targeting protein has been displayed on the surface of a minicell protoplast, these protoplasts are ready to be exposed to target cells. For preliminary experiments these interactions will be monitored using cell culture of Cos cells in comparison to lipofectamine (Invitrogen), electroporation, and other transfection techniques. Initial experiments will expose protoplasts displaying T-Inv to Cos cells and compare the transfection efficiency to protoplast containing pMPX-6::t-inv in the absence of t-inv expression, naked pMPX-6::t-inv alone, and naked pMPX-6::t-inv with lipofectamine. Each of these events will be monitored using fluorescent microscopy and/or flow cytometry. From these results the specific targeting apparatus proteins will be tested. Using A-431 (display EGFR) and K-562 (no EGFR) cell lines, the pMPX-6::t-egf constructs will be tested. Using the same approaches as for the t-inv study, the level of transfection between A-431 and K-562 cell lines will be measured and compared to those achieved using lipofectamine. Similarly, the ability of TNF to stimulate gene transfer will be studied using L-929 cells. In all cases, the ability of these general and specific targeting protein constructs will be compared to standard transfection techniques. Upon positive results, these methodologies will be tested on difficult to transfect cell lines, e.g. adult cardiomyocytes. The basis of these results will create a foundation for which applications into in vivo gene transfer may occur.

#### Example 20

##### Additional and Optimized Methods for Genetic Expression

[1201] Expression in minicells may occur following purification of minicells and/or minicell protoplasts from parental cells and LPS constituents, respectively. However, for some applications it is preferred to co-express proteins of interest with minicell induction. For these approaches, one may use the protocol described in Example 13 for expression of the *phoA* constructs. Either of these approaches may be accomplished using one or more of the following expression constructs (Table 23) and/or optimized expression constructs (Table 25).

[1202] Expression plasmid pCGV1 contains a temperature sensitive lambda cI repressor (cI857) and both lambda PR and PL promoters (Guzman, C. A., et al. 1994. A novel *Escherichia coli* expression-export vector containing alkaline phosphatase as an insertional inactivation screening system. *Gene.* 148:171-172) with an *atpE* initiation region (Schauder, B., et al. 1987. Inducible expression vectors incorporating the *Escherichia coli* *atpE* translational initiation region. *Gene.* 52:279-283). Included in the design of the

invention is the modification of this expression vector to best align the required Shine-Delgarno ribosomal binding site with cloning sites. In addition, the pCGVI expression vector was modified to incorporate a stem-loop structure at the 3-prime end of the transcript in order to provide a strong transcriptional stop sequence (Table 23).

[1203] Expression plasmid pCL478 contains a temperature sensitive lambda cI repressor (c1857) and both lambda PR and PL promoters (Love, C. A., et al. 1996. Stable high-copy bacteriophage promoter vectors for overproduction of proteins in *Escherichia coli*. Gene. 176:49-53). Included in the design of the invention is the modification of this expression vector to best align the required Shine-Delgarno ribosomal binding site with cloning sites. In addition, the pCL478 expression vector was modified to incorporate a stem-loop structure at the 3-prime end of the transcript in order to provide a strong transcriptional stop sequence (Table 23).

[1205] Oligonucleoides SEQ ID NOS.: 143 and 144 were annealed to each other to generate a DNA molecule with a 5' overhang at both ends. The overhangs are designed so that the DNA can be directly cloned into pCGVI cut with NdeI (5' overhang is TA) and BamHI (5' overhang is GATC). Insertion of the annealed DNA into pCGVI creates SEQ ID NO.: 139, pMPX-84.

[1206] Oligonucleoides SEQ ID NOS.: 145 and 146 were annealed to each other to generate a DNA molecule with a 5' overhang at both ends. The overhangs are designed so that the DNA can be directly cloned into pCGVI cut with NdeI (5' overhang is TA) and BamHI (5' overhang is GATC). Insertion of the annealed DNA into pCGVI creates SEQ ID NO.: 140, pMPX-85.

[1207] Oligonucleoides SEQ ID NOS.: 147 and 148 were annealed to each other to generate a DNA molecule with a 5' overhang at both ends. The overhangs are designed so that the DNA can be directly cloned into pCL478 cut with BamHI (5' overlap is GATC) and XhoI (overhang is TCGA).

TABLE 23

LAMBDA C1857 EXPRESSION VECTOR MODIFICATIONS				
New Plasmid	Parent plasmid	Region removed	Region added <sup>a</sup>	SEQ ID NO
pMPX-84	pCGVI	NdeI-BamHI	NdeI, SD-PstI, XbaI, KpnI, Stem-loop, BamHI	139
pMPX-85	pCGVI	NdeI-BamHI	NdeI, SD-SalI, XbaI, KpnI, Stem-loop, BamHI	140
pMPX-86	pCL478	BamHI-XhoI	BamHI, SD-PstI, XbaI, KpnI, Stem-loop, XhoI	141
pMPX-87	pCL478	BamHI-XhoI	BamHI, SD-SalI, XbaI, KpnI, Stem-loop, XhoI	142

<sup>a</sup> "SD" refers to a Shine-Delgarno ribosome-binding sequence; "Stem-loop" refers to a stem-loop structure that functions as a transcriptional stop site.

[1204]

TABLE 24

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 23		
SEQ ID NO	Primer name	5' to 3' sequence
143	CGV1-1-SalI	TATGTAAGGAGGTTGTCGACCGGCTCAGTCTAGAGGTACCCGCCCTCA TCCGAAAGGGCGTATTG
144	CGV1-2-SalI	GATCCAATACGCCCTTTCGGATGAGGGCGGGTACCTCTAGACTGAGCC GGTCGACAACCTCCTTACA
145	CGV1-1-PstI	TATGTAAGGAGGTTCTGCAGCGGCTCAGTCTAGAGGTACCCGCCCTCA TCCGAAAGGGCGTATTG
146	CGV1-2-PstI	GATCCAATACGCCCTTTCGGATGAGGGCGGGTACCTCTAGACTGAGCC GCTGCAGAACCTCCTTACA
147	CL478-1-SalI	GATCCTAAGGAGGTTGTCGACCGGCTCAGTCTAGAGGTACCCGCCCTC ATCCGAAAGGGCGTATTG
148	CL478-2-SalI	TCGAGAATACGCCCTTTCGGATGAGGGCGGGTACCTCTAGACTGAGCC GGTCGACAACCTCCTTAG
149	CL478-1-PstI	GATCCTAAGGAGGTTCTGCAGCGGCTCAGTCTAGAGGTACCCGCCCTC ATCCGAAAGGGCGTATTG
150	CL478-2-PstI	TCGAGAATACGCCCTTTCGGATGAGGGCGGGTACCTCTAGACTGAGCC GCTGCAGAACCTCCTTAG



Insertion of the annealed DNA into pCL578 cut with BamHI and XhoI creates SEQ ID NO.: 141, pMPX-86.

[1208] Oligonucleotides SEQ ID NOS.: 149 and 150 were annealed to each other to generate a DNA molecule with a 5' overhang at both ends. The overhangs are designed so that the DNA can be directly cloned into pCL578 cut with BamHI (5' overlap is GATC) and XhoI (overhang is TCGA). Insertion of the annealed DNA into pCL478 cut with BamHI and XhoI creates SEQ ID NO.: 142, pMPX-87.

[1209] The optimized expression constructs in Table 25 were created from SEQ ID NOS.: 6, 11, and 12 (see Table 13). Modifications were made to optimize the alignment of

TABLE 25

EXPRESSION CONSTRUCTS				
Plasmid	Regulatory element(s)	inducer	Plasmid	SEQ ID NO.:
pMPX-67	RhaRS	Rhamnose	PUC-18	151
pMPX-72	RhaRS	Rhamnose	PUC-18	152
pMPX-66	AraC	Arabinose	PUC-18	153
pMPX-71	AraC	Arabinose	PUC-18	154
pMPX-68	MelR	Melibiose	PUC-18	155

[1210]

TABLE 26

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 25 CONSTRUCTS		
SEQ ID NO.:	Primer name	5' to 3' sequence
69	Rha-1	GCGAATTGAGATGACGCCACTGGC
156	Rha-SD	GCAGAACCTCCTGAATTTTCATTACGACC
71	Rha-1-HindIII	CGGCGAAGCTTAATTAATCTTTCTGCGAATTGAGATGACGCCACTGGC
157	Rha-SD SalI KpnI	CCGCGGGTACCAATACGCCCTTTCGGATGAGGGCGCGGGG ATCCTCTAGAGTCGACGTCGACAACCTCCTGAATTTTCATTA CGACC
158	Rha-SD KpnI KpnI	CCGCGGGTACCAATACGCCCTTTCGGATGAGGGCGCGGGG ATCCTCTAGAGTCGACCTGCAGAACCTCCTGAATTTTCATTA CGACC
81	Ara-1	CAAGCCGTCAATTGTCTGATTTCG
159	Ara-SD	CTGCAGGGCCTCCTGTAGCCCCAAAAACGGGTATGG
83	Ara-1-HindIII	GCGCCAAGCTTCAAGCCGTCAATTGTCTGATTTCG
160	Ara-SD SalI KpnI	CCGCGGGTACCAATACGCCCTTTCGGATGAGGGCGCGGGG ATCCTCTAGAGTCGACGTCGACGGCCTCCTGTAGCCCCAAA AAAACGGGTATGG
161	Ara-SD PstI KpnI	CCGCGGGTACCAATACGCCCTTTCGGATGAGGGCGCGGGG ATCCTCTAGAGTCGACCTGCAGGGCCTCCTGTAGCCCCAAA AAAACGGGTATGG
77	Mel-1	CGTCTTTAGCCGGGAAACG
162	Mel-SD	CCTCCTGGCTTGCTTGAATAACTTCATCATGG
79	Mel-1-HindIII	GCCGCAAGCTTCGTCTTTAGCCGGGAAACG
163	Mel-SD-SalI KpnI	CCGCGGGTACCAATACGCCCTTTCGGATGAGGGCGCGGGG ATCCTCTAGAGTCGACCCCTCCTGGCTTGCCTGAATAACT TCATCATGGC

the SalI or PstI cloning sites with the Shine-Delgarno ribosome-binding site. In addition, stem-loop transcriptional termination sequences were added on the 3' end of the cloning region.

[1211] Oligonucleotides SEQ ID NOS.: 69, 156, 72, and 157 were used to amplify the rhaRS genes and their divergent control region from the *E. coli* chromosome and insertion of an optimized SalI-Shine-Delgarno ribosome--

binding alignment and a stem-loop transcriptional termination sequence. Once amplified, this region was inserted into pUC18 using HindIII and KpnI to create pMPX67, SEQ ID NO.: 151.

[1212] Oligonucleotides SEQ ID NOS.: 69, 156, 72, and 158 were used to amplify the rhaRS genes and their divergent control region from the *E. coli* chromosome and insertion of an optimized PstI-Shine-Delgarno ribosome-binding alignment and a stem-loop transcriptional termination sequence. Once amplified, this region was inserted into pUC18 using HindIII and KpnI to create, pMPX-72, SEQ ID NO.: 152.

[1213] Oligonucleotides SEQ ID NOS.: 81, 159, 81, 160 were used to amplify the araC genes and their divergent control region from the *E. coli* chromosome and insertion of an optimized SalI-Shine-Delgarno ribosome-binding alignment and a stem-loop transcriptional termination sequence. Once amplified, this region was inserted into pUC18 using HindIII and KpnI to create, pMPX-66, SEQ ID NO.: 153.

[1214] Oligonucleotides SEQ ID NOS.: 81, 159, 81, 161 were used to amplify the araC genes and their divergent control region from the *E. coli* chromosome and insertion of an optimized PstI-Shine-Delgarno ribosome-binding alignment and a stem-loop transcriptional termination sequence.

Once amplified, this region was inserted into pUC18 using HindIII and KpnI to create pMPX-71, SEQ ID NO.: 154.

[1215] Oligonucleotides SEQ ID NOS.: 77, 162, 79, 163 were used to amplify the meIR genes and their divergent control region from the *E. coli* chromosome and insertion of an optimized SalI-Shine-Delgarno ribosome-binding alignment and a stem-loop transcriptional termination sequence. Once amplified, this region was inserted into pUC18 using HindIII and KpnI to create, pMPX-68, SEQ ID NO.: 155.

#### Example 21

##### Optimization of Rat Neurotensin Receptor (NTR) Expression

[1216] Expression of specific GPCR proteins in minicells may require chimeric domain fusions to stabilize the expressed protein and/or direct the synthesized protein to the membrane. The NTR protein from rat was cloned into several chimeric combinations to assist in NTR expression and membrane association (Grisshammer, R., et al. 1993. Expression of a rat neurotensin receptor in *Escherichia coli*. *Biochem. J.* 295:571-576; Tucker, J., and Grisshammer, R. 1996. Purification of a rat neurotensin receptor expressed in *Escherichia coli*. *Biochem. J.* 317:891-899). Methods for construction are shown the Tables below.

TABLE 27

NEUROTENSIN RECEPTOR EXPRESSION FACILITATING CONSTRUCTS		
Protein <sup>a</sup>	Construct <sup>b</sup>	SEQ ID NO
MalE(L)	SalI-MalE (1-370)-Factor Xa-NTR homology	164
NTR	Factor Xa-NTR (43-424)-NotI-FLAG-KpnI	165
MalE(L)-NTR	SalI-MalE(1-370)-Factor Xa-NTR(43-424)-NotI-FLAG-KpnI	166
MalE(S)-NTR	SalI-MalE(1-28)-Factor Xa-NTR(43-424)-NotI-FLAG-KpnI	167
TrxA	NotI-TrxA(2-109)-NotI	168
MalE(L)-NTR-TrxA	SalI-MalE(1-370)-Factor Xa-NTR(43-424)-NotI-TrxA(2-109)-FLAG-KpnI	169
MalE(S)-NTR-TrxA	SalI-MalE(1-28)-Factor Xa-NTR(43-424)-NotI-TrxA(2-109)-FLAG-KpnI	170

<sup>a</sup>(L) refers to MalE residues 1-370, and (S) refers to MalE residues 1-28.

<sup>b</sup>All mature constructs were cloned into SalI and KpnI sites of SEQ ID NOS.: 140, 142, 151 and 153.

#### [1217]

TABLE 28

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 27		
SEQ ID NO.:	Primer name	5' to 3' sequence
171	MalE-1	GGTGCACGCATCCTCGCATTATCCGC
172	MalE-2	CGCACGGCATAACCAGAAAGCGGACATCTGCC
173	MalE-1-SalI	CCGCGGTTCGACATGAAAATAAAAAACAGGTGCACGCATCCTCGC
174	MalE-2-XaNTR	GCCGTGTCCGATTCGGAGGTGCGGCCTTCGATACGCACGGCAT ACCAAGAAAGCGGGATGTTCCGGC

TABLE 28-continued

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 27		
SEQ ID NO.:	Primer name	5' to 3' sequence
175	NTR-1	CCTCGGAATCCGACACGGCAGGGC
176	NTR-2	GTACAGGGTCTCCGGGTGGCGCTGG
177	NTR-1-Xa	CCGCGATCGAAGGCCGCACCTCGGAATCCGACACGGCAGGGCC
178	NTR-2-Flag	GGCGGGTACCTTTGTTCATCGTCATCTTTATAATCTGCGGCCGC GTACAGGGTCTCCGGGTGGCGCTGGTG
179	NTR-2-Stop KpnI	GGCGGGTACCTTATTATTGTTCATCGTCATCTTTATAATCTGC GGCCGC
180	NTR-1-Xa Lead	CCGCATTAACGACGATGATGTTTTCCGCCTCGGCTCTGCCAAA ATCATCGAAGGCCGCACCTCGGAATCCGACACGGC
181	NTR-2-Lead2 SalI	CCGCGGTCGACATGAAAATAAAAACAGGTGCACGCATCCTCGC ATTATCCGCATTAACGACGATGATGTTTTCCGCCTCGGC
182	TrxA-1	CCGCGAGCGATAAAATTATTACCTGACTGACG
183	TrxA-2	GCCCGCCAGGTTAGCGTCGAGGAACCTTTTCAACTGACC
184	TrxA-1-NotI	GCGGCCGCAAGCGATAAAATTATTACCTGACTGACG
185	TrxA-2-NotI	GGCGCTGCGCCGCATCATCATGATCTTTATAATCGCC

[1218] Oligonucleotides SEQ ID NOS.: 171, 172, 173 and 174 were used to amplify malE residues 1-370 from the *E. coli* chromosome to create SEQ ID NO.: 164. Using overlap PCR with the extended NTR homology, a chimeric translational fusion was made between MalE (1-370) and NTR residues 43-424 (SEQ ID NO.: 165) to create a SEQ ID NO.: 166. SEQ ID NO.: 166 was cloned into plasmids pMPX-85, pMPX-87, pMPX-66 and pMPX-67 (respectively, SEQ ID NOS.: 140, 142, 151 and 153) using SalI and KpnI.

[1219] Three-step PCR with oligonucleotides, SEQ ID NOS.: 175 and 176 as primers was used to amplify NTR residues 43-424 from rat brain cDNA. SEQ ID NOS.: 177 and 178 were then used with the NTR (43-424) template to add factor Xa and FLAG sequence. Finally, SEQ ID NOS.: 177 and 179 were used to add a KpnI site to create SEQ ID NO.: 165. Using overlap PCR with malE (1-370) containing extended NTR homology, a chimeric translational fusion was made between NTR (43-424) and MalE (1-370) (SEQ ID NO.: 164) to create a SEQ ID NO.: 166. SEQ ID NO.: 166 was cloned into SEQ ID NOS.: 140, 142, 151 and 153 using SalI and KpnI.

[1220] Using three-step PCR oligonucleotides SEQ ID NOS.: 175 and 176 were first used to amplify NTR residues 43-424 from rat brain cDNA. SEQ ID NOS.: 178 and 180 were then used with the NTR (43-424) template to add factor Xa and FLAG sequence. Finally, SEQ ID NOS.: 179 and 181 were used to add KpnI to create SEQ ID NO.: 167. SEQ ID NO.: 167 was cloned into SEQ ID NOS.: 140, 142, 151 and 153 using SalI and KpnI.

[1221] Oligonucleotides SEQ ID NOS.: 182, 183, 184 and 185 were used to amplify TrxA residues 2-109 from the *E. coli* chromosome to create SEQ ID NO.: 168. Using NotI, TrxA residues 2-109 was cloned into SEQ ID NOS.: 166 and 167 to create SEQ ID NOS.: 169 and 170, respectively. SEQ ID NOS.: 169 and 170 were cloned into SEQ ID NOS.: 140, 142, 151 and 153 using SalI and KpnI.

#### Example 22

##### Methods for Functional GPCR Assay

[1222] Functional G-protein-coupled receptor (GPCR) binding assays in minicells requires expression of a GPCR of interest into the minicell membrane bilayer and cytoplasmic expression of the required G-protein. For these purposes, constructs were created to co-express both a GPCR and a G-protein. To regulate the ratio of GPCR to G-protein, transcriptional fusions were created. In these constructs, the GPCR and G-protein are co-transcribed as a bi-cistronic mRNA. To measure the GPCR-G-protein interaction in the intact minicell, each protein was created as a chimera with a transactivation domain. For these studies the N-terminal DNA-binding, activation domain of the ToxR protein from *V. cholerae* was fused to the C-terminus of both the GPCR and G-protein. Finally, to measure the interaction GPCR-G-protein interaction, the ToxR-activated ctx promoter region was cloned in front of lacZ. Dimerization of the ToxR DNA-binding region will bind and activate the ctx promoter. In this construct, heterodimerization of the GPCR and G-protein will promote dimerization of ToxR that will be monitored by LacZ expression. Details of these constructs are shown in Table 29.

TABLE 29

FUNCTIONAL HUMAN GPCR CONSTRUCTS		
Protein <sup>a, b</sup>	Construct <sup>a, b</sup>	SEQ ID NO.:
$\beta$ 2AR	SalI- $\beta$ 2AR-PstI, XhoI	186
GS1 $\alpha$	XhoI-GS1 $\alpha$ -XbaI	187
$\beta$ 2AR-GS1 $\alpha$ fusion	SalI- $\beta$ 2AR-PstI, XhoI-GS1 $\alpha$ XbaI	188
$\beta$ 2AR-stop	SalI- $\beta$ 2AR-PstI-Stop-SD-XhoI	189
$\beta$ 2AR-stop-GS1 $\alpha$	SalI- $\beta$ 2AR-PstI-Stop-SD-XhoI-GS1 $\alpha$ XbaI	190
ToxR	ClaI-ToxR-XbaI	191
GS1 $\alpha$	XhoI-GS1 $\alpha$ -ClaI	192
GS2 $\alpha$	XhoI-GS2 $\alpha$ -ClaI	193
Gaq	XhoI-Gq $\alpha$ -ClaI	194
Gi $\alpha$	XhoI-Gi $\alpha$ -ClaI	195
G $\alpha$ 12/13	XhoI-G $\alpha$ 12/13-ClaI	196
GS1 $\alpha$ -ToxR	XhoI-GS1 $\alpha$ -ClaI-ToxR-XbaI	197
GS2 $\alpha$ -ToxR	XhoI-GS2 $\alpha$ -ClaI-ToxR-XbaI	198
Gaq-ToxR	XhoI-Gaq-ClaI-ToxR-XbaI	199
Gi $\alpha$ -ToxR	XhoI-Gi $\alpha$ -ClaI-ToxR-XbaI	200
G $\alpha$ 12/13-ToxR	XhoI-G $\alpha$ 12/13-ClaI-ToxR-XbaI	201
ToxR	PstI-ToxR-XhoI	202
$\beta$ 2AR	SalI- $\beta$ 2AR-PstI	203
$\beta$ 2AR-ToxR	SalI- $\beta$ 2AR-PstI-ToxR-Stop-SD-XhoI	204
$\beta$ 2AR-ToxR-stop-	SalI- $\beta$ 2AR-PstI-ToxR-Stop-SD-XhoI-GS1 $\alpha$ -ClaI-ToxR-XbaI	205
GS1 $\alpha$ -ToxR		
Pctx	XbaI-Pctx-lacZ homology	206
lacZ	Pctx homology-lacZ-XbaI	207
Pctx::lacZ	XbaI-Pctx-lacZ-XbaI	208

<sup>a</sup> "SD" refers to the Shine-Delgarno ribosome-binding sequence and "ToxR" refers to the transactivation, DNA-binding domain of the ToxR protein (residues 5-141).

<sup>b</sup> All mature constructs were cloned into SalI and XbaI sites of SEQ ID NOS.: 140, 142, 151 and 153.

[1223]

TABLE 30

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 29.		
SEQ ID NO.:	Primer name	5' to 3' sequence
209	$\beta$ 2AR-1	GGGGCAACCCGGGACGGCAGCGCC
210	$\beta$ 2AR-2	GCAGTGAGTCATTTGTACTACAATTCCTCC
211	$\beta$ 2AR-1-SalI	CGCGGTCGACATGGGGCAACCCGGGACGGCAGCGCC

TABLE 30-continued

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 29.		
SEQ ID NO.:	Primer name	5' to 3' sequence
212	$\beta$ 2AR-2-Link- XhoI	GGCTCGAGCTGCAGGTTGGTGACCGTCTGGCCACGCTC TAGCAGTGAGTCATTTGTACTACAATTCC
213	GS1 $\alpha$ -1	GGGCTGCCTCGGGAACAGTAAGACCGAGG
214	GS1 $\alpha$ -2	GAGCAGCTCGTACTGACGAAGGTGCATGC
215	GS1 $\alpha$ -1-XhoI	GGAGGCCCTCGAGATGGGCTGCCTCGGGAACAGTAAG ACCGAGG
216	GS1 $\alpha$ -2-XbaI	CCTCTAGATTATTATCGATGAGCAGCTCGTACTGACGA AGGTGCATGC
217	GS1 $\alpha$ -2-ClaI	CCATCGATGAGCAGCTCGTACTGACGAAGGTGCATGC
218	G $\alpha$ 12-1	CCGGGGTGGTGCGGACCCTCAGCCGC
219	G $\alpha$ 12-2	CTGCAGCATGATGTCCTTCAGGTTCTCC
220	G $\alpha$ 12-1-XhoI	GCGGGCTCGAGATGTCGGGGTGGTGCGGACCCTCAGC CGC
221	G $\alpha$ 12-2-ClaI	GCGCCATCGATCTGCAGCATGATGTCCTTCAGGTTCTCC
222	G $\alpha$ q-1	GACTCTGGAGTCCATCATGGCGTGTCTGC
223	G $\alpha$ q-2	CCAGATTGTACTCCTTCAGGTTCAACTGG
224	G $\alpha$ q-1-XhoI	ATGACTCTGGAGTCCATCATGGCGTGTCTGC
225	G $\alpha$ q-2-ClaI	GCGCCATCGATGACCAGATTGTACTCCTTCAGGTTCAACTGG
226	G $\alpha$ i-1	GGGCTGCACCGTGAGCGCCGAGGACAAGG
227	G $\alpha$ i-2	CCTTCAGGTTGTTCTTGATGATGACATCGG
228	G $\alpha$ i-1-XhoI	ATGGGCTGCACCGTGAGCGCCGAGGACAAGG
229	G $\alpha$ i-2-ClaI	GCGCCATCGATGAAGAGGCCGCGAGTCTTCAGGTTGTTCTTGA TGATGACATCGG
230	GS2 $\alpha$ -1	GGGCTGCCTCGGGAACAGTAAGACCGAGG
231	GS2 $\alpha$ -2	GAGCAGCTCGTACTGACGAAGGTGCATGC
232	GS2 $\alpha$ -1-XhoI	ATGGGCTGCCTCGGGAACAGTAAGACCGAGG
233	GS2 $\alpha$ -2-ClaI	GCGCCATCGATGAGCAGCTCGTACTGACGAAGGTGCATGC
234	$\beta$ 2AR-2-Link- Stop-XhoI	GGCTCGAGGGCCTCCTTGATTATTACTCGAGGGCCTCC TTGATTATTACTGCAGGTTGGTGACCGTCTGGCCACGC TCTAGCAGTGAGTCATTTGTACTACAATTCC
235	$\beta$ 2AR-2-Link	CCCTGCAGGTTGGTGACCGTCTGGCCACGCTCTAGCAG TGAGTCATTTGTACTACAATTCC
236	Tox (5-141)-1B	GGACACAAC TAAAAGAGATATCGATGAGTCATATTG G
237	Tox (5-141)-2	GAGATGTCATGAGCAGCTTCGTTTTTCGCG
238	Tox (5-141)-1- Link	CGGTGGCCAGACGGTCACCAACTGCAGGGACACAAC TAAAAGAGATATCG
239	Tox (5-141)-2- XhoI	CGGGATCCTCTAGATTATTAAGAGATGTCATGAGCAG CTTCGTTTTTCGCG
240	Ctx-1	GGCTGTGGGTAGAAGTGAAACGGGGTTTACCG
241	Ctx-2	CTTTACCATATAATGCTCCCTTGTTTTAAACAG

TABLE 30-continued

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 29.		
SEQ ID NO.:	Primer name	5' to 3' sequence
242	Ctx-2-XbaI	CGCGGTCTAGAGGCTGTGGGTAGAAGTGAACGGGGT TTACCG
243	Ctx-2-LacZ	CGACGGCCAGTGAATCCGTAATCATGGTCTTTACCATA TAATGCTCCCTTTGTTTAACAG
244	LacZ-1	CCATGATTACGGATTCACTGGCCGTCG
245	LacZ-2	CCAGACCAACTGGTAATGGTAGCGACC
246	LacZ-1-Ctx	GGTAAAGACCATGATTACGGATTCACTGGCCGTCG
247	LacZ-2-XbaI	GCGCCTCTAGAAATACGCCCTTTCGGATGAGGGCGTT ATTATTTTTGACACCAGACCAACTGGTAATGGTAGCG ACC

[1224] Oligonucleotides SEQ ID NOS.: 209, 210, 211 and 212 were used to amplify human  $\beta$ 2AR from human cDNA to create SEQ ID NO.: 186. Using Sall and XhoI a translational fusion was made between  $\beta$ 2AR and human GS1 $\alpha$  (SEQ ID NO.: 187) to create a SEQ ID NO.: 188. SEQ ID NO.: 188 was cloned into SEQ ID NOS.: 140, 142, 151 and 153 using Sall and XbaI.

[1225] Oligonucleotides SEQ ID NOS.: 213, 214, 215 and 216 were used to amplify human GS1 $\alpha$  from human cDNA to create SEQ ID NO.: 187. Using XhoI and XbaI a translational fusion was made between GS1 $\alpha$  and human  $\beta$ 2AR (SEQ ID NO.: 186) create SEQ ID NO.: 188. SEQ ID NO.: 188 was cloned into SEQ ID NOS.: 140, 142, 151 and 153 using Sall and XbaI.

[1226] Oligonucleotides SEQ ID NOS.: 213, 214, 215 and 217 were used to amplify human GS1 $\alpha$  from human cDNA to create SEQ ID NO.: 192. Using XhoI and XbaI a translational fusion was made with ToxR residues 5-141 from *Vibrio cholerae* (SEQ ID NO.: 191) to create SEQ ID NO.: 197. To be used to create a transcriptional fusion with  $\beta$ 2AR-ToxR chimeras as shown in SEQ ID NO.: 205 and future GPCR-ToxR chimeras.

[1227] Oligonucleotides SEQ ID NOS.: 218, 219, 220 and 221 were used to amplify human G $\alpha$ 12/13 from human cDNA to create SEQ ID NO.: 196. Using XhoI and XbaI a translational fusion was made h ToxR residues 5-141 from *Vibrio cholerae* (SEQ ID NO.: 191) to create SEQ ID NO.: 201. To be used to create future transcriptional fusions with GPCR-ToxR chimeras as shown in SEQ ID NO.: 205.

[1228] Oligonucleotides SEQ ID NOS.: 222, 223, 224 and 225 were used to amplify human G $\alpha$ q from human cDNA to create SEQ ID NO.: 194. Using XhoI and XbaI a translational fusion was made with ToxR residues 5-141 from *Vibrio cholerae* (SEQ ID NO.: 191) to create SEQ ID NO.: 199. To be used to create future transcriptional fusions with GPCR-ToxR chimeras as shown in SEQ ID NO.: 205.

[1229] Oligonucleotides SEQ ID NOS.: 226, 227, 228 and 229 were used to amplify human G $\alpha$ i from human cDNA to create SEQ ID NO.: 195. Using XhoI and XbaI a translational fusion was made with ToxR residues 5-141 from

*Vibrio cholerae* (SEQ ID NO.: 191) to create SEQ ID NO.: 200. To be used to create future transcriptional fusions with GPCR-ToxR chimeras as shown in SEQ ID NO.: 205.

[1230] Oligonucleotides SEQ ID NOS.: 230, 231, 232 and 233 were used to amplify human GS2 $\alpha$  from human cDNA to create SEQ ID NO.: 193. Using XhoI and XbaI a translational fusion was made with ToxR residues 5-141 from *Vibrio cholerae* (SEQ ID NO.: 191) to create SEQ ID NO.: 198. To be used to create future transcriptional fusions with GPCR-ToxR chimeras as shown in SEQ ID NO.: 205.

[1231] Oligonucleotides SEQ ID NOS.: 209, 210, 211 and 234 were used to amplify human P2AR from human cDNA to create SEQ ID NO.: 189. Using Sall and XhoI a transcriptional fusion was made between  $\beta$ 2AR and human GS1 $\alpha$  (SEQ ID NO.: 187) to create a SEQ ID NO.: 190. SEQ ID NO.: 190 was cloned into SEQ ID NOS.: 140, 142, 151 and 153 using Sall and XbaI.

[1232] Oligonucleotides SEQ ID NOS.: 236, 237, 238 and 239 were used to amplify bases coinciding with ToxR residues 5-141 from *Vibrio Cholerae* to create SEQ ID NO.: 202. Using PstI and XhoI a translational fusion was made between ToxR and human  $\beta$ 2AR (SEQ ID NO.: 203) to create SEQ ID NO.: 204.

[1233] Oligonucleotides SEQ ID NOS.: 209, 210, 211 and 235 were used to amplify human  $\alpha$ 2AR from human cDNA to create SEQ ID NO.: 203. Using Sall and PstI a translational fusion was made between  $\beta$ 2AR and ToxR (SEQ ID NO.: 202) to create SEQ ID NO.: 204.

[1234] Using oligonucleotides SEQ ID NOS.: 197 and 204 transcriptional fusions were created between the  $\beta$ 2AR-ToxR translational fusion (SEQ ID NO.: 204) and the GS1 $\alpha$ -ToxR translational fusion (SEQ ID NO.: 197) to create SEQ ID NO.: 205.

[1235] Oligonucleotides SEQ ID NOS.: 240, 241, 242 and 243 were used to amplify the ctx promoter region (Pctx) from *Vibrio cholerae* to create SEQ ID NO.: 206. Combining this PCR product in combination with the SEQ ID NO.: 207 PCR product and amplifying in the presence of SEQ ID NOS.: 242, 247, SEQ ID NO.: 208 was created. Using XbaI,

the SEQ ID NO.: 208 reporter construct was subsequently cloned into pACYC184 for co-transformation with the GPCR-G-protein fusions constructs above.

[1236] Oligonucleotides SEQ ID NOS.: 244, 245, 246 and 247 were used to amplify the lacZ from *E. coli* to create SEQ ID NO.: 207. Combining this PCR product in combination with the SEQ ID NO.: 206 PCR product and amplifying in the presence of SEQ ID NOS.: 242 and 247, SEQ ID NO.: 208 was created. Using XbaI, the 208 reporter construct was subsequently cloned into pACYC184 for co-transformation with the GPCR-G-protein fusions constructs above.

### Example 23

#### Modular Membrane-Targeting and Solubilization Expression Constructs

[1237] To produce membrane proteins efficiently in mini-cells it may be necessary to create chimeric fusions with the

membrane protein of interest. In this Example various regions of the MalE protein have been cloned into a modular expression system designed to create chimeric fusions with direct difficult to target membrane proteins to produce leader domains that will direct the proteins to the cytoplasmic membrane (Miller, K., W., et al. 1998. Production of active chimeric pediocin AcH in *Escherichia coli* in the absence of processing and secretion genes from the *Pediococcus* pap operon. *Appl. Environ. Microbiol.* 64:14-20). Similarly, a modified version of the TrxA protein has also been cloned into this modular expression system to create chimeric fusions with proteins that are difficult to maintain in a soluble conformation (LaVallie, E. R., et al. 1993. A thioredoxin gene fusion expression system that circumvents inclusion body formation in the *E. coli* cytoplasm. *Biotechnology* (N.Y.) 11:187-193). Table 31 describes each of these modular constructs.

TABLE 31

MODULAR MEMBRANE-TARGETING AND SOLUBILIZATION EXPRESSION CONSTRUCTS		
Protein <sup>a</sup>	Construct <sup>a</sup>	SEQ ID NO
MalE (1-28)	NsiI-MalE(1-28)-Factor Xa-PstI, SalI, XbaI-FLAG, NheI	248
MalE (1-370, del 354-364)	NsiI-MalE(1-370, del 354-364)-Factor Xa-PstI, SalI, XbaI-FLAG, NheI	249
TrxA (2-109, del 103-107)	PstI, SalI, XbaI-TrxA(2-109, del 103-107)-FLAG-NheI	250
MalE (1-28)-TrxA (2-109, del 103-107)	NsiI-MalE(1-28)-Factor Xa-PstI, SalI, XbaI-TrxA (2-109 del 103-107)-FLAG, NheI	251
MalE (1-370, del 354-364)-TrxA (2-109, del 103-107)	NsiI-MalE(1-370, del 354-364)-Factor Xa-PstI, SalI, XbaI-TrxA (2-109 del 103-107)-FLAG, NheI	252

<sup>a</sup>The term "del" refers to a deletion in which amino acid residues following the term "del" are removed from the sequence.

[1238]

TABLE 32

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 31.		
SEQ ID NO.:	Primer name	5' to 3' sequence
253	MalE-1-NsiI	CGCGGATGCATATGAAAATAAAAAACAGGTGCACGCATCCTCGCATT ATCCGCATTAACGACGATGATGTTTTCCGCCTCGGCTCTCGCC
254	MalE-2-middle	CGTCGACCGAGGCTGCAGGCGGCTTCGATGATTTTGCGCAG AGCCGAGGCGGAAAACATCATCGTCG
255	MalE-3s-NheI	CGAAGCCCGCTGCAGGCTCGGTCGACGCCGAATCTAGAGATTAT AAAGATGACGATGACAAAATAATAAGCTAGCGGCGC
256	MalE-4-NheI	GCGCCGCTAGCTTATTATTTGTCATCG
257	MalE-1a	GGTGCACGCATCCTCGCATTATCCGC
258	MalE-2a	GGCGTTTTCCATGGTGGCGGCAATACGTGG

TABLE 32-continued

OLIGONUCLEOTIDE PRIMER SEQUENCES FOR TABLE 31.		
259	MalE-1-NsiI	CGCGGATGCATATGAAAATAAAAAACAGGTGCACGCATCCTC GCATTATCCGC
260	MalE-2-NheI	CCGAGGCCTGCAGGCGGGCTTCGATACGCACGGCATAACCAG AAAGCGGACTGGCGTTTTCATGGTGGCGCAATACGTGG
261	MalE-3L-NheI	GCGCCGCTAGCTTATTATTTGTCATCGTCATCTTTATAATCTC TAGATTTCGGCGTCGACCGAGGCTGCAGGCGGGCTTCGATA CGC
262	TrxA-1a	CCTGACTGACGACAGTTTTGACACGG
263	TrxA-2a	CCTTTAGACAGTGACCCACTTTGGTTGCCGC
264	TrxA-1a-PsiI	CGCGGCTGCAGGCCTCGGTGCACGCCGAATCTAGAAGCGAT AAAATTATTCACCTGACTGACGACAGTTTTGACACGG
265	TrxA-2-NheI	GCGCCGCTAGCTTATTATTTGTCATCGTCATCTTTATAATCCG CCAGGTTCTCTTTCAACTGACCTTTAGACAGTGACCCACTTT GGTTGCCGC

[1239] Oligonucleotides SEQ ID NOS.: 253, 254, 255 and 256 overlap with each other to form a scaffold template to PCR amplify malE (1-28) to create a SEQ ID NO.: 248. Following PCR amplification, SEQ ID NO.: 248 was digested with NsiI and NheI and cloned into SEQ ID NOS.: 152, 154, 139 and 141 digested with PstI and XbaI. The resultant products create SEQ ID NOS.: 266, 267, 268 and 269, respectively, that lose both the 5-prime PstI and 3-prime XbaI restriction site and retain the PstI, SalI, and XbaI restriction sites between MalE (1-28) and the FLAG sequence. Insertion of a protein in alignment with these sites results in a chimeric protein containing amino-terminal MalE (1-28) and carboxy-terminal FLAG.

[1240] Oligonucleotides SEQ ID NOS.: 257, 258, 259 and 260 were used to amplify malE (1-370 with a deletion removing residues 354-364) to create SEQ ID NO.: 249. Following PCR amplification, SEQ ID NO.: 249 was digested with NsiI and NheI and cloned into SEQ ID NOS.: 152, 154, 139 and 141 digested with PstI and XbaI. The resultant products create SEQ ID NOS.: 270, 271, 272 and 273, respectively, that lose both the 5-prime PstI and 3-prime XbaI restriction site and retain the PstI, SalI, and XbaI restriction sites between MalE (1-370, del 354-364) and the FLAG sequence. Insertion of a protein in alignment with these sites results in a chimeric protein containing amino-terminal MalE (1-370, del 354-364) and carboxy-terminal FLAG.

[1241] Oligonucleotides SEQ ID NOS.: 262, 263, 264 and 265 were used to amplify trxA (2-109 with a deletion removing residues 103-107) to create SEQ ID NO.: 250. Following PCR amplification, SEQ ID NO.: 250 was digested with PstI and NheI and cloned into SEQ ID NOS.: 152, 154, 139 and 141 digested with PstI and XbaI. to create SEQ ID NOS.: 274, 275, 276 and 277, respectively. Using these restriction digestion combinations results in loss of the XbaI SEQ ID NO.: 249 insertion site.

[1242] The resultant products create SEQ ID NOS.: 274, 275, 276 and 277, respectively, that lose the 3-prime XbaI restriction site and retain the PstI, SalI, and XbaI restriction

sites on the 3-prime end of the TrxA (1-109, del 103-107) sequence. Insertion of a protein in alignment with these sites results in a chimeric protein containing Carboxy-terminal TrxA (1-109, del 103-107)-FLAG.

[1243] SEQ ID NO.: 248 was digested with NsiI and XbaI and cloned into SEQ ID NOS.: 274, 275, 276 and 277 that were digested with PstI and XbaI. The resultant products create SEQ ID NOS.: 278, 279, 280 and 281, respectively, that lose the 5 prime PstI restriction site and retain the PstI, SalI, and XbaI restriction sites between MalE (1-28) and TrxA (1-109, del 103-107). Insertion of a protein in alignment with these sites results in a chimeric protein containing amino-terminal MalE (1-28) and carboxy-terminal TrxA (1-109, del 103-107)-FLAG.

[1244] SEQ ID NO.: 249 was digested with NsiI and XbaI and cloned into SEQ ID NOS.: 274, 275, 276 and 277 that were digested with PstI and XbaI. The resultant products create SEQ ID NOS.: 282, 283, 284 and 285, respectively, that lose the 5 prime PstI restriction site and retain the PstI, SalI, and XbaI restriction sites between MalE (1-370, del 354-364) and TrxA (1-109, del 103-107). Insertion of a protein in alignment with these sites results in a chimeric protein containing amino-terminal MalE (1-370, del 354-364) and carboxy-terminal TrxA (1-109, del 103-107)-FLAG.

#### Example 24

##### Poroplast Formation

[1245] Minicells are used to prepare Poroplasts in order to increase the accessibility of a membrane protein component and/or domain to the outside environment. Membrane proteins in the inner membrane are accessible for ligand binding and/or other interactions in poroplasts, due to the absence of an outer membrane. The removal of the outer membrane from *E. coli* whole cells and minicells to produce poroplasts was carried out using modifications of previously described protoplast and analysis protocols (Birdsell et al., Production and Ultrastructure of Lysozyme and Ethylenediaminetet-



raacetate-Lysozyme Spheroplasts of *Escherichia coli*, J. Bacteriol. 93:427-437, 1967; Weiss et al., Protoplast Formation in *Escherichia Coli*, J. Bacteriol. 128:668-670, 1976; Matsuyama, S-I., et al. SecD is involved in the release of translocated secretory proteins from the cytoplasmic membrane of *Escherichia coli*. 12:265-270, 1993).

[1246] In brief, cells were grown to late-log phase and pelleted at room temperature. Minicells were also isolated from cultures in late-log phase. The pellet was washed twice with 50 mM Tris, pH 8.0. Following the second wash,  $1 \times 10^9$  cells were resuspended in 1 ml 50 mM Tris (pH 8.0) that contained 8% sucrose and 2 mM EDTA. Cell/EDTA/sucrose mixtures were incubated at 37° C. for 10 min, centrifuged, decanted, and poroplasted cells were resuspended in 50 mM Tris, pH 8.0 with 8% sucrose. Incubation with anti-LPS-coated magnetic beads, as described in Example 14, is used to enrich for poroplasts that lack LPS. Following incubation with the resuspended protoplasted cells, the anti-LPS magnetic beads were removed from suspension with a magnet.

[1247] To examine the range of molecular sizes that can pass through the cell wall, an IgG molecule was tested for its ability to pass the intact cell wall. Binding of an antibody to the ToxR-PhoA chimera expressed on the inner membrane minicell poroplasts was measured. Briefly, minicell poroplasts with and without inner membrane-bound ToxR-PhoA were incubated at 37° C. with anti-PhoA antibody in reaction buffer (50 mM Tris, pH 8.0, 8% sucrose, 1% BSA, and 0.01% Tween-20). Following incubation, poroplasts were centrifuged, washed 3 times with reaction buffer, and resuspended in 50 mM Tris, pH 8.0 with 8% sucrose. Following resuspension, bound proteins from  $5 \times 10^7$  minicells or minicell poroplasts were separated using denaturing SDS-PAGE, transferred to nitrocellulose, and developed using with both anti-PhoA antibody and secondary antibody against both heavy and light chains of anti-PhoA IgG (Table 33).

TABLE 33

ANTI-PHOA ACCESSIBILITY TO POROPLAST INNER MEMBRANE-BOUND TOXR-PHOA				
EDTA (mM)	0	2	0	2
Lysozyme (mg/ml)	0	0	5	5
	Poroplasts (ng antibody bound)		Protoplasts (ng antibody bound)	
Minicells ToxR- PhoA	ND <sup>a</sup>	0.6	ND <sup>a</sup>	12.8
Minicells only	ND <sup>a</sup>	ND <sup>a</sup>	ND <sup>a</sup>	ND <sup>a</sup>

<sup>a</sup>Non-detectable

[1248] These results demonstrate that the cell wall present on poroplasts is penetrable by an IgG molecule and that an IgG molecule is capable of passing the intact cell wall and binding to an inner membrane protein. From this data it appears that poroplast operate at ~10% the efficiency of protoplasts by allowing 0.6 ng of IgG to bind inner membrane-bound ToxR-PhoA compared to 12.8 ng. However, given the large size of IgG (~150,000 Daltons) it is expected that molecules having a smaller molecular weight will efficiently access inner membrane proteins in poroplasts.

## Example 25

## Production of Neurotensin Receptor (NTR)

[1249] To demonstrate expression of NTR in isolated minicells, MalE(L)-NTR (SEQ ID NO.: 166 was cloned into pMPX-67 (SEQ ID NO.: 151). Following minicell isolation,  $1.5 \times 10^9$  minicells were induced with 1 mM Rhamnose for 2 hour at 37° C. Following induction, the protein produced was visualized via Western analysis using an anti-MalE antibody following separation on an SDS-PAGE. The results are shown in FIG. 2.

[1250] These data demonstrates that MalE(L)-NTR is induced 87-fold by addition of 1 mM rhamnose to the minicell induction mixture. Cross-reactive proteins are host MalE and non-specific binding by Goat-anti-mouse HRP secondary antibody.

[1251] The contents of the articles, patents, and patent applications, and all other documents and electronically available information mentioned or cited herein, are hereby incorporated by reference in their entirety to the same extent as if each individual publication was specifically and individually indicated to be incorporated by reference. Applicants reserve the right to physically incorporate into this application any and all materials and information from any such articles, patents, patent applications, or other documents.

[1252] The inventions illustratively described herein may suitably be practiced in the absence of any element or elements, limitation or limitations, not specifically disclosed herein. Thus, for example, the terms "comprising", "including," containing", etc. shall be read expansively and without limitation. Additionally, the terms and expressions employed herein have been used as terms of description and not of limitation, and there is no intention in the use of such terms and expressions of excluding any equivalents of the features shown and described or portions thereof, but it is recognized that various modifications are possible within the scope of the invention claimed. Thus, it should be understood that although the present invention has been specifically disclosed by preferred embodiments and optional features, modification and variation of the inventions embodied therein disclosed may be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of this invention.

[1253] The invention has been described broadly and generically herein. Each of the narrower species and sub-generic groupings falling within the generic disclosure also form part of the invention. This includes the generic description of the invention with a proviso or negative limitation removing any subject matter from the genus, regardless of whether or not the excised material is specifically recited herein.

[1254] Other embodiments are within the following-claims. In addition, where features or aspects of the invention are described in terms of Markush groups, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group.

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 SEQUENCE LISTING
 

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&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

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&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

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gctgattaat	aatctagagg	cgttaccaat	tatgacaact	tgacgggaag	ttcctattct	2220
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&lt;210&gt; SEQ ID NO 6

&lt;211&gt; LENGTH: 4728

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 6

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cagcttgtct	gtaagcggat	gccgggagca	gacaagcccg	tcagggcgcg	tcagcgggtg	120
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gattcgcggt	attgcagaaa	gccatcccgt	ccttggcgaa	tatcacgcgg	tgaccagtta	1380
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taggggttcc gcgcacattt ccccgaaaag tgccacctga cgtctaagaa accattatta 4680
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<210> SEQ ID NO 7
<211> LENGTH: 1440
<212> TYPE: DNA
<213> ORGANISM: E. coli

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<400> SEQUENCE: 7

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tttaacgggt gatcagactg ccgctctcgt tgattctctt agcgataaac ctgcaaaaaa 180
tattattttg ctgattggcg atgggatggg ggactcggaa attactgccg cacgtaatta 240
tgccgaaggt gcgggcgctt tttttaaagg tatagatgcc ttaccgctta ccgggcaata 300
cactcactat gcgtgtaata aaaaaaccgg caaacggac tacgtcaccg actcggctgc 360
atcagcaacc gcctgtgcaa ccggtgtcaa aacctataac ggcgcgctgg gcgtcgatat 420
tcacgaaaaa gatcacccaa cgattctgga aatggcaaaa gccgcaggtc tggcgaccgg 480
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gccagtgcgc tggctaggac cgaagaacaac gtaccatggc aatatacgata agcccgcagt	900
cacctgtagc ccaaatccgc aacgtaatac cagtgtacca accctggcgc agatgaccga	960
caaagccatt gaattgttga gtaaaaatga gaaaggcttt ttctgcaag ttgaagggtc	1020
gtcaatcgat aaacaggatc atgctgcaaa tccttggggg caaattggcg agacggtcga	1080
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gggcctcacc caggcgttaa ataccaaaaga tggcgagctg atgggatgga gttacgggaa	1260
ctccgaagag gattcacaag aacataccgc cagtcagttg cgtattgagg cgtatggccc	1320
gcatgccgcg aatgtttgtg gactgaccga ccagaccgat ctcttctaca ccatgaaagc	1380
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<210> SEQ ID NO 8  
 <211> LENGTH: 1560  
 <212> TYPE: DNA  
 <213> ORGANISM: E. coli

<400> SEQUENCE: 8

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aaataaagt aaacaaagca ctattgcact ggcactctta ccgttactgt ttaccctgt	180
gacaaaagcc cggacaccag aaatgcctgt tctggaaaac cgggctgctc agggcgatat	240
tactgcaccc ggcgggtgctc gccgtttaac ggggtgatcag actgccgctc tgcgtgattc	300
ttctagcgat aaacctgcaa aaaatattat tttgctgatt ggcgatggga tgggggactc	360
ggaaattact gccgcacgta attatgccga aggtgcgggc ggcTTTTTTT aaggtataga	420
tgcttaccg cttaccgggc aatacactca ctatgcgctg aataaaaaaa ccggcaaacc	480
ggactacgct accgactcgg ctgcatcagc aaccgcctgg tcaaccgggt tcaaaacctc	540
taacggcgcg ctgggcgctg atattcacga aaaagatcac ccaacgattc tggaaatggc	600
aaaagccgca ggtctggcga ccggtaacgt ttctaccgca gagttgcagg atgccacgcc	660
cgctgcgctg tgggcacatg tgacctcgcg caaatgctac ggtccgagcg cgaccagtga	720
aaaatgtccg ggtaacgctc tggaaaaagg cggaaaaagg tcgattaccg aacagctgct	780
taacgctcgt gccgacgta cgcttggcgg cggcgcaaaa acctttgctg aaacggcaac	840
cgctggtgaa tggcagggaa aaacgctcgc tgaacaggca caggcgcgctg gttatcagtt	900
ggtgagcgat gctgcctcac tgaattcggg gacggaaagc aatcagcaaa aacctctgct	960
tggcctgttt gctgacggca atatgccagt gcgctggcta ggaccgaaag caacgtacca	1020
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tgttgcgccc	gataccaaag	ctccgggccc	caccaggcgc	ctaaatacca	aagatggcgc	1380
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gttgcgattt	gcggcgatg	gcccgcagtc	cgccaatggt	gttgactga	ccgaccagac	1500
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<210> SEQ ID NO 9  
 <211> LENGTH: 1500  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 9

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<210> SEQ ID NO 10  
 <211> LENGTH: 3908  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Expression vector

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caggagctca attaatacac taacggaccg gtaacaacc gtgcgtgttg tttaccggga	480
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ggataacatg cagactgaag cacaaccgac acggaccggg atcctcaatg ctgccagaga	720
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&lt;210&gt; SEQ ID NO 11

&lt;211&gt; LENGTH: 3872

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 11

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ttggcgggtg tcggggctgg ctttaactatg cggcatcaga gcagattgta ctgagagtg 180
accatagcgc ggtgaaata ccgcacagat gcgtaaggag aaaataccgc atcagggcgc 240

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&lt;210&gt; SEQ ID NO 12

&lt;211&gt; LENGTH: 3934

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 12

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thggcgggtg thcggggctgg ctthactatg cggcatcaga gcagattgth ctgagagthc	180
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gcacggaact cgtcgggct ggcccgggtg cththththaa athaccgcga gaaatagagth	540
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&lt;210&gt; SEQ ID NO 13

&lt;211&gt; LENGTH: 5953

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 13

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<210> SEQ ID NO 14  
 <211> LENGTH: 1380  
 <212> TYPE: DNA  
 <213> ORGANISM: Rat

<400> SEQUENCE: 14

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cacctcatc accaccatcc tcttcttggg cacctgtagc ttcatcgtct tggagaacct	240
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cggcaacttg gctctctgcg acctgctggc cggcatagcc tacaaggta acattctgat	360
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<210> SEQ ID NO 15  
 <211> LENGTH: 1320  
 <212> TYPE: DNA

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&lt;213&gt; ORGANISM: Homo sapien

&lt;400&gt; SEQUENCE: 15

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gccggaccac gacgtcacgc agcaaaggga cgagggtggt gtgggtgggca tgggcatcgt      180
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caagttcgag cgtctgcaga cggtcaccaa ctacttcata acttcactgg cctgtgctga      300
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caagtaccag agcctgctga ccaagaataa ggcccgggtg atcattctga tgggtgtggat      540
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&lt;210&gt; SEQ ID NO 16

&lt;211&gt; LENGTH: 52

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Cloning primer

&lt;400&gt; SEQUENCE: 16

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aattggtacc tcaatgatga tgatgatgat gcttcagag gacccattc tg                52

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&lt;210&gt; SEQ ID NO 17

&lt;211&gt; LENGTH: 1320

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Homo sapien

&lt;400&gt; SEQUENCE: 17

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tcgcaactct ctactgtttc tccataccgg ttttttggg ctagcaggag gaattcacca      60
tggatagtgt gtgtcccaa ggaaatata tccacctca aaataattcg atttgotgta     120
ccaagtgcc caaaggaacc tacttgtaata atgactgtcc aggcccggg caggatacgg     180
actgcaggga gtgtgagagc ggctccttca cgccttcaga aaaccacctc agacactgcc     240

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tgctgggacg cgtgctccgc gacatggacc tgctgggctg cctggaggac atcagaggag	1260
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&lt;210&gt; SEQ ID NO 18

&lt;211&gt; LENGTH: 1380

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Homo sapien

&lt;400&gt; SEQUENCE: 18

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tccctcacct aggggacagg gagaagagag atagtgtgtg tccccaagga aaatataatc	120
accctcaaaa taattcgatt tgctgtacca agtgccaaa aggaacctac ttgtacaatg	180
actgtccagg cccggggcag gatacggact gcagggagtg tgagagcggc tccctcaccg	240
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gtttctttct aagagaaaa gagtgtgtct cctgtagtaa ctgtaagaaa agcctggagt	540
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cagtgtgttt gccctggtc attttctttg gtctttgcct tttatccctc ctcttcattg	660
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<210> SEQ ID NO 19
<211> LENGTH: 780
<212> TYPE: DNA
<213> ORGANISM: Homo sapien

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<400> SEQUENCE: 19

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gagcccatct atctgggagg ggtcttccag ctggagaagg gtgaccgact cagcgotgag 660
atcaatcggc ccgactatct cgactttgcc gagtctgggc aggtctactt tgggatcatt 720
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<210> SEQ ID NO 20
<211> LENGTH: 300
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Gene encoding a fusion protein

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<400> SEQUENCE: 20

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ttattctgat agcggctctta cttcccctcg cagtattact gctcaatagt gactctgaat 120
gtcccctgtc ccacgatggg tactgcctcc atgatggtgt gtgcatgtat attgaagcat 180
tggacaagta tgcattgcaac tgtgttgttg gctacatcgg ggagcgatgt cagtaccgag 240
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<210> SEQ ID NO 21
<211> LENGTH: 1620
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Gene encoding a fusion protein

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&lt;400&gt; SEQUENCE: 21

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ttgagtttta ctcaaacgg tgtgccggtg agtattagcc ccattaccga gcagccagat	900
agctataccg cgacggtggt tgggaatagt gtcggtgatg tcacaatcac gccgcaggtt	960
gataccctga tactgagtac attgcagaaa aaaatatccc tattcccggg acctacgctg	1020
accggtattc tggttaacgg gcaaaatttc gctacggata aagggttccc gaaaacgac	1080
tttaaaaacg ccacattcca gttacagatg gataacgat ttgctaataa tactcagtat	1140
gagtggctgt cgtcattcac acccaatgta tcgggtaacg atcaggggtca ggtgacgatt	1200
acctacaaa cctatagcga agtggtgtg acggcgaaaa gtaaaaaatt cccaagtatt	1260
tcggtgagtt atcggttcta cccaaatcgg tggatatacg atggcggcag atcgttggt	1320
tccagtctcg aggcagcag acaatgcca ggttcagata tgtctgcggt tcttgaatcc	1380
tcacgtgcaa caacggaac gcgtgcgcct gacgggacat tgtggggcga gtgggggagc	1440
ttgaccgcgt atagttctga ttggcaatct ggtgaatatt gggtaaaaa gaccagcacg	1500
gattttgaaa ccatgaatat ggacacagc gcaactgcaac cagggcctgc atacttggcg	1560
ttcccgcctc gtgcgctgtc aatataactg caggcatgca agcttggccc gcgggcccgg	1620

&lt;210&gt; SEQ ID NO 22

&lt;211&gt; LENGTH: 208

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: E. coli

&lt;400&gt; SEQUENCE: 22

gaattcaggc gctttttaga ctggtcgtaa tgaaattcag caggatcaca ttctgcagat	60
gtcacggcgc agacttatag tcgctttgtt tttatTTTTT aatgtatttg tacatggaga	120
aaataaagtg aaacaaagca ctattgcact ggcaacttta cgttactgt ttaccctgt	180
gacaaaagcc cggacaccag aatctaga	208

&lt;210&gt; SEQ ID NO 23

&lt;211&gt; LENGTH: 1546

&lt;212&gt; TYPE: DNA



-continued

&lt;213&gt; ORGANISM: E. coli

&lt;400&gt; SEQUENCE: 23

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gaattcaggc gctttttaga ctggtcgtaa tgaattcag caggatcaca ttctgcagat    60
gtcacggcgc agacttatag tcgctttggt tttatttttt aatgtatttg tacatggaga    120
aaataaagtg aaacaaagca ctattgcaact ggcaactctta cgttactgt ttaccctgt    180
gacaaaagcc cggacaccag aaatgcctgt tctggaaaac cgggctgctc agggcgatat    240
tactgcaccc ggcggtgctc gccgtttaac gggatgatcag actgccgctc tgcgtgattc    300
tcttagcgat aaacctgcaa aaaatattat tttgctgatt ggcgatggga tgggggactc    360
ggaaattact gccgcacgta attatgccga aggtgcgggc ggctttttta aaggtataga    420
tgccttaccc cttaccgggc aatacactca ctatgcgctg aataaaaaaa ccggcaaac    480
ggactacgtc accgactcgg ctgcatcagc aaccgcctgg tcaaccggtg tcaaacctta    540
taacggcgcg ctgggcctgc atattcacga aaaagatcac ccaacgattc tggaaatggc    600
aaaaaccgca ggtctggcga ccggtaacgt ttctaccgca gatttgacag atgccacgcc    660
cgctgcgctg gtggcacatg tgacctcgcg caaatgtac ggtccgagcg cgaccagtga    720
aaaaatgtcc ggtaacgctc tggaaaaagg cggaaaaaga tcgattaccg aacagctgct    780
taacgctcgt gccgacgtta cgcttgccgg cggcgcaaaa acctttgctg aaacggcaac    840
cgctggtgaa tggcagggaa aaacgctgcg tgaacaggca caggcgcgtg gttatcagtt    900
ggtgagcgat gctgcctcac tgaattcggg gacggaaagc aatcagcaaa aaccctgct    960
tggcctgttt gctgacggca atatgccagt gcgctggcta ggaccgaaag caacgtacca   1020
tggcaatata gataagcccg cagtcacctg tacgccaaat ccgcaacgta atgacagtgt   1080
accaaccctg gcgcagatga ccgacaaaag cattgaattg ttgagtaaaa atgagaaaag   1140
ctttttcctg caagttgaag gtgcgtcaat cgataaacag gatcatgctg cgaatccttg   1200
tgggcaaatg ggcgagacgg tcgatctcga tgaagccgta caacggggcg tggaaatcgc   1260
taaaaaggag ggtaacacgc tggatcatag caccgctgat caccgccacg ccagccagat   1320
tgttgccgcg gataccaaa gctccggcct caccaggcg ctaaatacca aagatggcgc   1380
agtgatggtg atgagttacg ggaactccga agaggattca caagaacata ccggcagtca   1440
gttgcgtatt gcggcgatg gcccgcatgc cgccaatggt gttggactga ccgaccagac   1500
cgatctcttc tacaccatga aagccgctct ggggctgaaa tctaga                    1546

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&lt;210&gt; SEQ ID NO 24

&lt;211&gt; LENGTH: 148

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: E. coli

&lt;400&gt; SEQUENCE: 24

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gaattcaggc gctttttaga ctggtcgtaa tgaattcag caggatcaca ttctgcagat    60
gaaaataaaa acaggtgcac gcatcctcgc attatccgca ttaacgacga tgatgttttc   120
cgcctcggct ctcgcaaaa tctctaga                    148

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&lt;210&gt; SEQ ID NO 25

&lt;211&gt; LENGTH: 1174

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: E. coli

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&lt;400&gt; SEQUENCE: 25

gaattcaggc gctttttaga ctggctgtaa tgaattcag caggatcaca ttctgcagat	60
gaaaataaaa acaggtgcac gcatcctcgc attatccgca ttaacgacga tgatgttttc	120
cgctcggct ctcgccaaaa tcgaagaag taaactggta atctggatta acggcgataa	180
aggctataac ggtctcgcctg aagtcggtaa gaaattcgag aaagataccg gaattaaagt	240
caccgttgag catccggata aactggaaga gaaattccca caggttgccg caactggcga	300
tggccctgac attatcttct gggcacacga cgcctttggt ggctacgctc aatctggcct	360
gttggtgtaa atcaccocgg aaaaagcgtt ccaggacaag ctgtatccgt ttacctggga	420
tgccgtacgt tacaacggca agctgattgc ttacccgatc gctgttgaag cgttatcgct	480
gatttataac aaagatctgc tgcgcaacc gccaaaaacc tgggaagaga tcccgccgct	540
ggataaagaa ctgaaagcga aaggtaaag cgcgctgatg ttcaacctgc aagaaccgta	600
cttcacctgg ccgctgattg ctgctgacgg gggttatgcy ttcaagtatg aaaacggcaa	660
gtacgacatt aaagacgtgg gcgtggataa cgctggcgcg aaagcgggtc tgaccttcct	720
ggttgacctg attaaaaaca aacacatgaa tgcagacacc gattactcca tcgcagaagc	780
tgctcttaat aaagcgaaa cagcgatgac catcaacggc ccgtgggcat ggtccaacat	840
cgacaccagc aaagtgaatt atggtgtaac ggtactgccg acctcaagg gtcaaccatc	900
caaaccgttc gttggcgtgc tgagcgcagg tattaacgcc gccagtccga acaagagct	960
ggcgaaagag ttctcgtgaa actatctgct gactgatgaa ggtctggaag cggttaataa	1020
agacaaacgc ctgggtgccg tagcgtgaa gtcttacgag gaagagtgg cgaaagatcc	1080
acgtattgcc gccaccatgg aaaacccca gaaaggtgaa atcatgccga acatcccga	1140
gatgtccgct ttctggtatg ccgtgcgttc taga	1174

&lt;210&gt; SEQ ID NO 26

&lt;211&gt; LENGTH: 3840

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 26

accatattgc ggtgaaata ccgcacagat gcgtaaggag aaaataccgc atcaggcgcc	60
atacgcgaca gcgcgcaata accgttctcg actcataaaa gtgatgccgc tataatgccg	120
cgctctatct gaatgctttc gggatgatc tggtaacagg gaatgtgatt gattataaga	180
acatcccggc tccgcgaagc caacaacctg tgcttgccgg gtaagagttg accgagcact	240
gtgatttttt gagtaacaa gatgcaagtt tcagttgaaa ccaactcaag ccttgccgc	300
cggtaacga ttactatcgc tgctgacagc atcgagaccg ctgttaaaag cgagctggtc	360
aacgttgcca aaaaagtacg tattgacggc ttccgcaaag gcaaagtgcc aatgaatatac	420
gttctcagc gttatggcgc gtctgtacgc caggacgttc tgggtgacct gatgagccgt	480
aacttcattg acgccatcat taaagaaaa atcaatccgg ctggcgcacc gacttatggt	540
ccggcgcaat acaagctggg tgaagacttc acttactctg tagagtttga agtttatccg	600
gaagttgaac tgcagggtct ggaagcgcgc gaagttgaaa aaccgatcgt tgaagtgacc	660
gacgctgaag ttgacggcat gctggatact ctgcgtaaac agcaggcgac ctggaaagaa	720
aaagacggcg ctgttgaagc agaagaccgc gtaaccatcg acttcaccgg ttctgtagac	780

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ggcgaagagt tcgaaggcgg taaagcgtct gatttcgtac tggcgatggg ccagggtcgt	840
atgatcccgg gctttgaaga cggatatcaa ggccacaaag ctggcgaaga gttcaccatc	900
gacgtgacct tcccgaaga ataccacgca gaaaacctga aaggtaaagc agcgaattc	960
gctatcaacc tgaagaaagt tgaagagcgt gaactgccgg aactgactgc agaattcatc	1020
aaacgtttcg gcgttgaaga tggttccgta gaaggctgc gcgctgaagt gcgtaaaaac	1080
atggagcggc agctgaagag cgcctaccgt aaccgcgta agtctcaggc gatcgaaggt	1140
ctggtaaaa ctaacgacat cgactaccg gctgcgctga tcgacagcga aatcgacttt	1200
ctgcgtcggc aggtgcgaca gcgtttcggg ggcaacgaaa aacaagctct ggaactgccg	1260
cgcaactgt tcgaagaaca ggctaaacgc cgcgtagttg ttggcctgct gctgggcgaa	1320
gttatccgca ccaacgagct gaaagctgac gaagagcggc tgaagggcct gatcgaagag	1380
atggcttctg cgtacgaaga tccgaagaa gttatcgagt tctacagcaa aaacaagaa	1440
ctgatggaca acatgcgcaa tgttgctctg gaagaacagg ctggtgaagc tgtactggcg	1500
aaagcgaag tgactgaaa agaaaccact ttcaacgagc tgatgaacca gcaggcgtaa	1560
taataatcta gaggtagcac aatcagattc gottatgacg gcgatgaaga aattgogatg	1620
aaatgtgag tgaatcaggg ttttcaccgg atttgtgct gatcagaatt tttttcttt	1680
ttccccctg aaggggcgaa gcctcatccc catttctctg gtcaccagcc gggaaaccac	1740
gtaagctcgg gcgtcaccca taacagatac ggactttctc aaaggagagt tatcaatgaa	1800
tattcgtcca ttgcatgac gcgtgatcgt caagcgtaaa gaagttgaaa ctaaatctgc	1860
tggcggcacc gttctgaccg gctctgcagc ggctaaatcc acccgcgggc aagtgtggc	1920
tgctcggcaat ggccgatcc ttgaaatgg cgaagtgaag ccgctggatg tgaagttgg	1980
cgacatcggt attttcaacg atggctacgg tgtgaaatct gagaagatcg acaatgaaga	2040
agtggtgatc atgtccgaaa gcgacattct ggcaattggt gaagcgtaat ccgcgcacga	2100
cactgaacat acgaatttaa ggaataaaga taatggcagc taaagacgta aaattcgta	2160
acgacgctcg tgtgaaaatg ctgcgcggcg taaacgtact ggacagatgca gtgaaagtta	2220
ccctcggtc aaaaggccgt aacgtagttc tggataaatc tttcggtgca ccgaccatca	2280
ccaaagatgg tgtttccggt gctcgtgaaa tcgaactgga agacaagttc gaaaatatgg	2340
gtgcgcagat ggtgaaagaa gttgcctcta aagcaaacga cgtgcaggc gacggtacca	2400
ccactgcaac cgtactggct caggctatca tcaactgaagg tctgaaagct gttgctcgg	2460
gcatgaacc gatggacctg aaacgtggta tcgacaaagc ggttaccgct gcagttgaag	2520
aactgaaagc gctgtccgta ccatgctctg actctaaagc gattgctcag gttggtacca	2580
tctccgctaa ctccgacgaa accgtaggta aactgatcgc tgaagcgatg gacaaagtcg	2640
gtaaagaag cgttatcacc gttgaagacg gtaccggtct gcaggacgaa ctggacgtgg	2700
ttgaaggtat gcagttcgac cgtggctacc tgtctcotta ctcatcaac aagccgaaa	2760
ctggcgcagt agaactggaa agcccgttca tctgctggc tgacaagaaa atctccaaca	2820
tccgcgaaat gctgcgggtt ctggaagctg ttgccaaagc aggcaaaccg ctgctgatca	2880
tcgctgaaga tgtagaagc gaagcgtgg caactctggt tgttaacacc atgctggca	2940
tcgtgaaagt cgtgcgggtt aaagcaccgg gottcggcga tcgtcgtaaa gctatgctgc	3000
aggatatacg aaccctgact ggcggtaccg tgatctctga agagatcggg atggagctgg	3060

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aaaaagcaac cctggaagac ctgggtcagg ctaaactgtg tgtgatcaac aaagacacca 3120
ccactatcat cgatggcgtg ggtgaagaag ctgcaatcca gggccgtgtt gctcagatcc 3180
gtcagcagat tgaagaagca acttctgact acgaccgtga aaaactgcag gaacgcgtag 3240
cgaaactggc aggcggcgtt gcagttatca aagtgggtgc tgctaccgaa gttgaaatga 3300
aagagaaaaa agcagcgcgtt gaagatgccc tgcacgcgac ccgtgctgcg gtagaagaag 3360
gcgtggttgc tgggtggtgt gttgcgctga tccgcgtagc gtctaaactg gctgacctgc 3420
gtggtcagaa cgaagaccag aacgtgggta tcaaagtgc actgctgca atggaagctc 3480
cgctgcgta gatcgtattg aactgcggcg aagaaccgtc tgttgttct aacacggtta 3540
aaggcggcga cggcaactac ggttacaacg cagcaaccga agaatacggc aacatgatcg 3600
acatgggtat cctggatcca accaaagtaa ctcgttctgc tctgcagtac gcagcttctg 3660
tggctggcct gatgatcacc accgaatgca tggttaccga cctgccgaaa aacgatgcag 3720
ctgacttagg cgctgctggc ggtatggcg gcattgggtg catggcgcg atgatgtaat 3780
aataagcttg catgcctgca ggtcgactct agaggatccc cgggtaccga gctcgaattc 3840

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<210> SEQ ID NO 27
<211> LENGTH: 480
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Gene encoding a fusion protein
<400> SEQUENCE: 27

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gaattcagcg cctttttaga ctggtcgtaa tgaaattcag caggatcaca ttctgcagat 60
gatcgaagcc cgctctagac tcgagagcga taaaattatt cacctgactg acgacagttt 120
tgacacggat gtactcaaa cggacggggc gatcctcgtc gatttctggg cagagtggtg 180
cggtcctgctg aaaaatgatc ccccgattct ggatgaaatc gctgacgaat atcagggcaa 240
actgaccggt gcaaaaactga acatcgatca aaacctggc actgcgccga aatatggcat 300
ccgtgggtat ccgactctgc tgctgttcaa aaacgggtgaa gtggcgccaa ccaaagtggg 360
tgactgtct aaaggtcagt tgaaagatt cctcgacgct aacctggcg tcgaggatta 420
taaagatcat gatggcgatt ataaagatca tgatgattaa taaggatccc cgggtaccga 480

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<210> SEQ ID NO 28
<211> LENGTH: 1140
<212> TYPE: DNA
<213> ORGANISM: Rat

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<400> SEQUENCE: 28

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atggcaacca cgcacgcgca ggggcaccgc ccagtcttgg ggaatgatac tctccgggaa 60
cattatgatt acgtggggaa gctggcagcg aggctgcggg atccccctga gggtagcacc 120
ctcatcacca ccacctctct cttggtcacc tgtagcttca tcgtcttga gaacctgatg 180
gttttgattg ccatctggaa aaacaataaa tttcataacc gcatgtactt tttcatcggc 240
aacttggctc tctgcgacct gctggccgca atagcctaca aggtcaacat tctgatgtcc 300
ggtaggaaga cgttcagcct gtctccaaca gtgtggttcc tcagggaggg cagtatgttc 360
gtagccctgg gcgcatccac atgcagctta ttggccattg ccattgagcg gcacctgacc 420
atgatcaaga tgaggccgta cgacgcaca aagaagcacc gcgtgttcct tctgattggg 480

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atgtgctggc taattgcctt ctgctgggt gccctgcca tcctgggctg gaactgcctg 540
gagaactttc cggactgctc taccatcttg cccctctact ccaagaaata cattgccttt 600
ctcatcagca tcttcacagc cattctgggt accatcgtca tcttgtagc gcgcatctac 660
ttcctggtoa agtccagcag ccgcagggtg gccaaccaaca actccgagag atccatggcc 720
cttctcgcca ccgtagtgat cgtggtgagc gtgttcatcg cctgttggtc cccccctttc 780
atcctcttcc tcatcgatgt ggctgcagg gcgaaggagt gctccatcct cttcaagagt 840
cagtggttca tcatgctggc tgtcctcaac tcggccatga accctgtcat ctacacgctg 900
gccagcaaaq agatgcggcg tgctttcttc cggttggtgt gcggctgtct ggtcaagggc 960
aaggggaccc aggcctcccc gatgcagcct gctcttgacc cgagcagaag taaatcaagc 1020
tccagtaaca acagcagcag ccaactctca aaggtcaagg aagacctgcc ccatgtggct 1080
acctcttctc gcgtcactga caaaacgagg tcgcttcaga atggggctct ctgcaagtga 1140

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&lt;210&gt; SEQ ID NO 29

&lt;211&gt; LENGTH: 379

&lt;212&gt; TYPE: PRT

&lt;213&gt; ORGANISM: Rat

&lt;400&gt; SEQUENCE: 29

```

Met Ala Thr Thr His Ala Gln Gly His Pro Pro Val Leu Gly Asn Asp
 1          5          10          15
Thr Leu Arg Glu His Tyr Asp Tyr Val Gly Lys Leu Ala Gly Arg Leu
          20          25          30
Arg Asp Pro Pro Glu Gly Ser Thr Leu Ile Thr Thr Ile Leu Phe Leu
          35          40          45
Val Thr Cys Ser Phe Ile Val Leu Glu Asn Leu Met Val Leu Ile Ala
          50          55          60
Ile Trp Lys Asn Asn Lys Phe His Asn Arg Met Tyr Phe Phe Ile Gly
          65          70          75          80
Asn Leu Ala Leu Cys Asp Leu Leu Ala Gly Ile Ala Tyr Lys Val Asn
          85          90          95
Ile Leu Met Ser Gly Arg Lys Thr Phe Ser Leu Ser Pro Thr Val Trp
          100          105          110
Phe Leu Arg Glu Gly Ser Met Phe Val Ala Leu Gly Ala Ser Thr Cys
          115          120          125
Ser Leu Leu Ala Ile Ala Ile Glu Arg His Leu Thr Met Ile Lys Met
          130          135          140
Arg Pro Tyr Asp Ala Asn Lys Lys His Arg Val Phe Leu Leu Ile Gly
          145          150          155          160
Met Cys Trp Leu Ile Ala Phe Ser Leu Gly Ala Leu Pro Ile Leu Gly
          165          170          175
Trp Asn Cys Leu Glu Asn Phe Pro Asp Cys Ser Thr Ile Leu Pro Leu
          180          185          190
Tyr Ser Lys Lys Tyr Ile Ala Phe Leu Ile Ser Ile Phe Thr Ala Ile
          195          200          205
Leu Val Thr Ile Val Ile Leu Tyr Ala Arg Ile Tyr Phe Leu Val Lys
          210          215          220
Ser Ser Ser Arg Arg Val Ala Asn His Asn Ser Glu Arg Ser Met Ala
          225          230          235          240
Leu Leu Arg Thr Val Val Ile Val Val Ser Val Phe Ile Ala Cys Trp

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	245		250		255										
Ser	Pro	Leu	Phe	Ile	Leu	Phe	Leu	Ile	Asp	Val	Ala	Cys	Arg	Ala	Lys
			260					265					270		
Glu	Cys	Ser	Ile	Leu	Phe	Lys	Ser	Gln	Trp	Phe	Ile	Met	Leu	Ala	Val
		275					280					285			
Leu	Asn	Ser	Ala	Met	Asn	Pro	Val	Ile	Tyr	Thr	Leu	Ala	Ser	Lys	Glu
	290					295					300				
Met	Arg	Arg	Ala	Phe	Phe	Arg	Leu	Val	Cys	Gly	Cys	Leu	Val	Lys	Gly
305				310						315					320
Lys	Gly	Thr	Gln	Ala	Ser	Pro	Met	Gln	Pro	Ala	Leu	Asp	Pro	Ser	Arg
				325					330					335	
Ser	Lys	Ser	Ser	Ser	Ser	Asn	Asn	Ser	Ser	Ser	His	Ser	Pro	Lys	Val
			340					345					350		
Lys	Glu	Asp	Leu	Pro	His	Val	Ala	Thr	Ser	Ser	Cys	Val	Thr	Asp	Lys
		355					360					365			
Thr	Arg	Ser	Leu	Gln	Asn	Gly	Val	Leu	Cys	Lys					
	370					375									

<210> SEQ ID NO 30  
 <211> LENGTH: 32  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 30

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<210> SEQ ID NO 31  
 <211> LENGTH: 32  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 31

aattggatcc ttaagaagaa gaattgacgt tt 32

<210> SEQ ID NO 32  
 <211> LENGTH: 32  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 32

aattggatcc agaagaagaa ttgacgtttc ca 32

<210> SEQ ID NO 33  
 <211> LENGTH: 52  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 33

aattggatcc ttaatgatga tgatgatgat gagaagaaga attgacgttt cc 52

<210> SEQ ID NO 34

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<211> LENGTH: 24  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 34  
  
ttatggcaac cacgcacgcg cagg 24  
  
<210> SEQ ID NO 35  
<211> LENGTH: 21  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 35  
  
agaccgtcac ttgcagagga c 21  
  
<210> SEQ ID NO 36  
<211> LENGTH: 32  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 36  
  
aattgctagc acgcacgcg aggggcacc gc 32  
  
<210> SEQ ID NO 37  
<211> LENGTH: 34  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 37  
  
aattggtacc tcacttgagc aggaccccat tctg 34  
  
<210> SEQ ID NO 38  
<211> LENGTH: 32  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 38  
  
aattgctagc acgcacgcg aggggcacc gc 32  
  
<210> SEQ ID NO 39  
<211> LENGTH: 21  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
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<223> OTHER INFORMATION: Cloning primer  
  
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<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 40

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<210> SEQ ID NO 41

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<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 41

ggtcgccacc atggtgagca a 21

<210> SEQ ID NO 42

<211> LENGTH: 31

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 42

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<211> LENGTH: 25

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 43

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<210> SEQ ID NO 44

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<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 44

gcttgcttac gcaggaatgc tggg 24

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<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 45

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<210> SEQ ID NO 46

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<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

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<400> SEQUENCE: 46



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<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 47

gctagactgg gcggttttat ggacagcaag c 31

<210> SEQ ID NO 48  
<211> LENGTH: 35  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 48

gcgttaataa ttcagaagaa ctgctcaaga aggcg 35

<210> SEQ ID NO 49  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 49

gcgctactg acgtagttcg acgctcggac tagcgaagtt cctatacttt ctagagaata 60

ggaacttcgc tagactgggc ggttttatgg acagcaagc 99

<210> SEQ ID NO 50  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 50

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agtataggaa cttcgcgtta ataattcaga agaactcgtc aagaaggcg 109

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<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 51

cgttaccaat tatgacaact tgacgg 26

<210> SEQ ID NO 52  
<211> LENGTH: 29  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 52

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tgacgg	66
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gatgacgcc	69
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tgtctttggt gaccag	76
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tgacgg	66
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<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 58

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<210> SEQ ID NO 59  
<211> LENGTH: 28  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 59

gcggctttca tgggttagaa gagatcgg 28

<210> SEQ ID NO 60  
<211> LENGTH: 43  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 60

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<210> SEQ ID NO 61  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 61

gcgcctctag attattattt cagccccaga gcggctttca tgggttagaa gagatcgg 58

<210> SEQ ID NO 62  
<211> LENGTH: 24  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 62

gtcacggcgg agacttatag tcgc 24

<210> SEQ ID NO 63  
<211> LENGTH: 28  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 63

gcggctttca tgggttagaa gagatcgg 28

<210> SEQ ID NO 64  
<211> LENGTH: 37  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 64

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ccgcgctgca gatgtcacgg ccgagactta tagtcgc 37

<210> SEQ ID NO 65  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 65

gcccctctag attattattt cagccccaga gggccttca tgggttagaa gagatcgg 58

<210> SEQ ID NO 66  
<211> LENGTH: 109  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 66

ccgcgctgca gatgaacttg ggaatcgc tgttattct gatagcggc ttacttccc 60

tcgcagtatt actgctcatg cctgttctgg aaaaccgggc tgctcagg 109

<210> SEQ ID NO 67  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 67

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<210> SEQ ID NO 68  
<211> LENGTH: 24  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 68

gcgaattgag atgacgccac tggc 24

<210> SEQ ID NO 69  
<211> LENGTH: 25  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 69

cctgctgaat ttcattaacg accag 25

<210> SEQ ID NO 70  
<211> LENGTH: 48  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 70

cggcgaagt taattaatct ttctgcaat tgagatgacg ccactggc 48

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<210> SEQ ID NO 71  
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<212> TYPE: DNA  
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<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 71  
  
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<210> SEQ ID NO 72  
<211> LENGTH: 22  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 72  
  
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<210> SEQ ID NO 73  
<211> LENGTH: 23  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 73  
  
cctcattaag ataataatac tgg 23

<210> SEQ ID NO 74  
<211> LENGTH: 33  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 74  
  
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<210> SEQ ID NO 75  
<211> LENGTH: 47  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 75  
  
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<210> SEQ ID NO 76  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 76  
  
cgtcttttagc cgggaaacg 19

<210> SEQ ID NO 77  
<211> LENGTH: 18  
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<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 77  
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<210> SEQ ID NO 78  
<211> LENGTH: 30  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
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<211> LENGTH: 26  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 79  
cggtcgacgc agatctcctg gcttgc 26  
  
<210> SEQ ID NO 80  
<211> LENGTH: 23  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 80  
caagccgtca attgtctgat tcg 23  
  
<210> SEQ ID NO 81  
<211> LENGTH: 22  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 81  
ggtgaattcc tcctgctagc cc 22  
  
<210> SEQ ID NO 82  
<211> LENGTH: 34  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 82  
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<210> SEQ ID NO 83  
<211> LENGTH: 28  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 83  
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<210> SEQ ID NO 84  
<211> LENGTH: 42  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 84  
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<210> SEQ ID NO 85  
<211> LENGTH: 34  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 85  
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<210> SEQ ID NO 86  
<211> LENGTH: 26  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 86  
ggcaaccacg cacgcgcagg gccacc 26

<210> SEQ ID NO 87  
<211> LENGTH: 25  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 87  
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<210> SEQ ID NO 88  
<211> LENGTH: 38  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 88  
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<210> SEQ ID NO 89  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
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<400> SEQUENCE: 89  
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<210> SEQ ID NO 90  
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<400> SEQUENCE: 90  
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<210> SEQ ID NO 91  
<211> LENGTH: 30  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
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gcagtgagtc attgtacta caattcctcc 30

<210> SEQ ID NO 92  
<211> LENGTH: 37  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
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cgcggtcgac atggggcaac cgggaacgg cagcgcc 37

<210> SEQ ID NO 93  
<211> LENGTH: 49  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 93  
gcgccggatc cttattatag cagtgagtc tttgtactac aattcctcc 49

<210> SEQ ID NO 94  
<211> LENGTH: 28  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 94  
ggactgtcc ctcacctagg ggacaggg 28

<210> SEQ ID NO 95  
<211> LENGTH: 27  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 95  
ctgagaagac tgggcgcggg cgggagg 27

<210> SEQ ID NO 96  
<211> LENGTH: 42  
<212> TYPE: DNA



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<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 96  
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<210> SEQ ID NO 97  
<211> LENGTH: 44  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 97  
gcgccgtac cttattactg agaagactgg ggcgggagg gagg 44  
  
<210> SEQ ID NO 98  
<211> LENGTH: 16  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 98  
gatagtgtgt gtcccc 16  
  
<210> SEQ ID NO 99  
<211> LENGTH: 17  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 99  
ctgagaagac tgggagc 17  
  
<210> SEQ ID NO 100  
<211> LENGTH: 27  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
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gggagacat ggatagtgtg tgtcccc 27  
  
<210> SEQ ID NO 101  
<211> LENGTH: 32  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 101  
gcctcatcta gattactgag aagactgggc gc 32  
  
<210> SEQ ID NO 102  
<211> LENGTH: 26  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 102

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<210> SEQ ID NO 103  
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 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 103

cagggcaatg atcccaaagt agacctgc 28

<210> SEQ ID NO 104  
 <211> LENGTH: 39  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 104

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<210> SEQ ID NO 105  
 <211> LENGTH: 43  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 105

ggcgcaagct tatcacaggg caatgatccc aaagtagacc tgc 43

<210> SEQ ID NO 106  
 <211> LENGTH: 110  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 106

tctgatagcg gtcttacttc cctctgcagt attactgctc aatagtgact ctgaatgtcc 60

cctgtcccac gatgggtact gcctccatga tgggtgtgtgc atgtatattg 110

<210> SEQ ID NO 107  
 <211> LENGTH: 101  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 107

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tccaatgctt caatatacat gcacacacca tcatggaggc a 101

<210> SEQ ID NO 108  
 <211> LENGTH: 63  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 108

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tcg 63

<210> SEQ ID NO 109

<211> LENGTH: 61

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 109

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<211> LENGTH: 22

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 110

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<210> SEQ ID NO 111

<211> LENGTH: 24

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 111

ttatattgac agcgcacaga gcgg 24

<210> SEQ ID NO 112

<211> LENGTH: 101

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 112

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<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 113

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<210> SEQ ID NO 114

<211> LENGTH: 24

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

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<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 114

gtcacggccg agacttatag tcgc 24

<210> SEQ ID NO 115  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 115

ggtgtccggg cttttgtcac agg 23

<210> SEQ ID NO 116  
<211> LENGTH: 37  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 116

cgcggtgca gatgtcacgg ccgagactta tagtcgc 37

<210> SEQ ID NO 117  
<211> LENGTH: 38  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 117

cgcggtctag attctggtgt ccgggctttt gtcacagg 38

<210> SEQ ID NO 118  
<211> LENGTH: 23  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 118

cagccccaga gcggctttca tgg 23

<210> SEQ ID NO 119  
<211> LENGTH: 37  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 119

cgcggtctag atttcagccc cagagcggct ttcattgg 37

<210> SEQ ID NO 120  
<211> LENGTH: 106  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 120

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cgatgatggt ttccgcctcg gctctcgcca aaatctctag acgcgg 106

<210> SEQ ID NO 121

<211> LENGTH: 106

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 121

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<210> SEQ ID NO 122

<211> LENGTH: 26

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 122

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<211> LENGTH: 26

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 123

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<210> SEQ ID NO 124

<211> LENGTH: 52

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 124

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<210> SEQ ID NO 125

<211> LENGTH: 42

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 125

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<210> SEQ ID NO 126

<211> LENGTH: 27

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 126

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cgcgacagcg cgcaataacc gttctcg 27

<210> SEQ ID NO 127  
<211> LENGTH: 27  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 127

gctggttcat cagctcgttg aaagtgg 27

<210> SEQ ID NO 128  
<211> LENGTH: 41  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 128

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<210> SEQ ID NO 129  
<211> LENGTH: 52  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 129

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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 130

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<211> LENGTH: 25  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 131

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<210> SEQ ID NO 132  
<211> LENGTH: 38  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
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<210> SEQ ID NO 133

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<211> LENGTH: 49  
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 <400> SEQUENCE: 133  
  
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<210> SEQ ID NO 134  
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 <212> TYPE: DNA  
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 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer  
  
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<210> SEQ ID NO 135  
 <211> LENGTH: 26  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer  
  
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<210> SEQ ID NO 136  
 <211> LENGTH: 66  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer  
  
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 ctgacg 66

<210> SEQ ID NO 137  
 <211> LENGTH: 101  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer  
  
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<210> SEQ ID NO 138  
 <211> LENGTH: 65  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer  
  
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<210> SEQ ID NO 139  
<211> LENGTH: 67  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 139

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<210> SEQ ID NO 140  
<211> LENGTH: 65  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 140

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<210> SEQ ID NO 141  
<211> LENGTH: 67  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 141

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<210> SEQ ID NO 142  
<211> LENGTH: 66  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 142

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<210> SEQ ID NO 143  
<211> LENGTH: 66  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 143

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<210> SEQ ID NO 144  
<211> LENGTH: 66  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:



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<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 144

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<210> SEQ ID NO 145

<211> LENGTH: 66

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 145

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<210> SEQ ID NO 146

<211> LENGTH: 4740

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Expression vector

<400> SEQUENCE: 146

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ttggcgggtg tcggggctgg cttactatg cggcatcaga gcagattgta ctgagagtgc	180
accatattgc gtgtgaaata ccgcacagat gcgtaaggag aaaataccgc atcaggcggc	240
attcggcatt caggctgcgc aactgttggg aagggcgatc ggtgcggggc tottcgctat	300
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tttcccagtc acgacgttgt aaaacgacgg ccagtgccaa gcttaattaa tctttctcgg	420
aattgagatg acgccactgg ctgggcgtca tcccggtttc ccgggtaaac accaccgaaa	480
aatagttaact atcttcaaa ccacattcgg tcgaaatata actgattaac aggcggctat	540
gctggagaag atattgcgca tgacacactc tgacctgtcg cagatattga ttgatgggca	600
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&lt;210&gt; SEQ ID NO 147

&lt;211&gt; LENGTH: 4746

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 147

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ttcgtc						4746

&lt;210&gt; SEQ ID NO 148

&lt;211&gt; LENGTH: 3946

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 148

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ttggcgggtg	tcggggctgg	cttaactatg	cggcatcaga	gcagattgta	ctgagagtgc	180
accatattgcg	gtgtgaaata	ccgcacagat	gcgtaaggag	aaaataccgc	atcagggccc	240
attcgccatt	caggctgcgc	aactgttggg	aagggcgatc	ggtgcgggcc	tcttcgctat	300

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tgattcgtta	ccaattatga	caacttgacg	gctacatcat	tcacttttcc	ttcacaaccg	480
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&lt;210&gt; SEQ ID NO 149

&lt;211&gt; LENGTH: 3952

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 149

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attcgcatt	caggctgcgc	aactgttggg	aagggcgatc	ggtgcgggcc	tcttcgctat	300
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gcacggaact	cgctcgggct	ggccccgggt	cattttttaa	atacccgcga	gaaatagagt	540
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&lt;210&gt; SEQ ID NO 150

&lt;211&gt; LENGTH: 3886

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Expression vector

&lt;400&gt; SEQUENCE: 150

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tcattcagct ccggttccca acgatcaagg cgagttacat gatcccccat gttgtgcaaa 3240
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cagggttatt gtctcatgag cggtacata tttgaatgta tttagaaaaa taaacaata 3780
ggggttcgcg gcacatttcc ccgaaaatg ccacctgacg tctaagaaac cattattatc 3840
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<210> SEQ ID NO 151
<211> LENGTH: 28
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 151
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gcagaacctc ctgaatttca ttacgacc 28
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<210> SEQ ID NO 152
<211> LENGTH: 86
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 152
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ccgcgggtac caatacgccc tttcggatga gggcgcgggg atcctctaga gtcgacgtcg 60
acaacctcct gaatttcatt acgacc 86

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<210> SEQ ID NO 153
<211> LENGTH: 86
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 153
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ccgcgggtac caatacgccc tttcggatga gggcgcgggg atcctctaga gtcgacctgc 60
agaacctcct gaatttcatt acgacc 86

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<210> SEQ ID NO 154
<211> LENGTH: 38
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 154
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ctgcagggcc tcctgctagc ccaaaaaaac gggtatgg 38
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<210> SEQ ID NO 155  
 <211> LENGTH: 94  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 155

ccgcgggtac caatacgccc ttctggatga gggcgcgggg atcctctaga gtcgacgtcg 60  
 acggcctcct gctagcccaa aaaaacgggt atgg 94

<210> SEQ ID NO 156  
 <211> LENGTH: 94  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 156

ccgcgggtac caatacgccc ttctggatga gggcgcgggg atcctctaga gtcgacctgc 60  
 agggcctcct gctagcccaa aaaaacgggt atgg 94

<210> SEQ ID NO 157  
 <211> LENGTH: 32  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 157

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<210> SEQ ID NO 158  
 <211> LENGTH: 91  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 158

ccgcgggtac caatacgccc ttctggatga gggcgcgggg atcctctaga gtcgaccccc 60  
 tcctggcttg cttgaataac ttcatcatgg c 91

<210> SEQ ID NO 159  
 <211> LENGTH: 2319  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 159

gtcgacatga aaataaaaac aggtgcacgc atcctcgcat tatccgcatt aacgacgatg 60  
 atgtttttccg cctcggtctc cgccaaaatc gaagaaggta aactggtaat ctggattaac 120  
 ggcgataaag gctataacgg tctcgctgaa gtcggtaaga aattcgagaa agataaccgga 180  
 attaaagtoa ccggtgagca tccggataaa ctggaagaga aattcccaca ggttgoggca 240  
 actggcggatg gccctgacat tatcttctgg gcacacgacc gctttgggtg ctacgctcaa 300  
 tctggcctgt tggctgaaat caccocggac aaagcgttcc aggacaagct gtatccggtt 360

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acctgggatg	ccgtacgtta	caacggcaag	ctgattgctt	accogatcgc	tgttgaagcg	420
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ccggcgctgg	ataaagaact	gaaagcgaaa	ggtaagagcg	cgctgatggt	caacctgcaa	540
gaacctgact	tcacctggcc	gctgattgct	gctgacgggg	gttatgcggt	caagtatgaa	600
aacggcaagt	acgacattaa	agacgtgggc	gtggataacg	ctggcgcgaa	agcgggtctg	660
accttcctgg	ttgacctgat	taaaaacaaa	cacatgaatg	cagacaccga	ttactocatc	720
gcagaagctg	cctttaataa	aggcgaaaca	gcgatgacca	tcaacggccc	gtgggcatgg	780
tccaacatcg	acaccagcaa	agtgaattat	ggtgtaacgg	tactgccgac	cttcaagggt	840
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aaagagctgg	cgaagaggtt	cctcgaaaac	tatctgctga	ctgatgaagg	tctggaagcg	960
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cgctacttgg	ccatctgcca	tccctcaag	gccaagacc	tcagtcccg	cagccgcacc	1560
aagaaattca	tcagtgccat	atggctagct	tggcgctgc	tggctatacc	catgcttttc	1620
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cccattgtgg	acacagccac	tgtcaaggtc	gtcatccagg	ttaacacctt	catgtccttc	1740
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atggtgcacc	aggccgcca	gcagggccga	gtgtgcaccg	tgggcacaca	caacggttta	1860
gagcacagca	cgttcaacat	gaccatcgag	ccgggtcgtg	tccaggccct	gcgccacgga	1920
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cgacgcctga	tgttctgcta	tatctcggat	gaacagtgga	ctacgttcct	cttcgatttc	2040
taccactatt	tctacatgct	aaccaacgct	ctcttctacg	tcagctccgc	catcaatccc	2100
atcctctaca	acctggtctc	cgccaacttc	cgccaggtct	ttctgtccac	gctggcctgc	2160
ctttgtcctg	ggtggcgcca	ccgccgaaa	aagaggccaa	cgttctccag	gaagcccaac	2220
agcatgtcca	gcaacctatg	cttttccacc	agcgcacccc	gggagaccct	gtacgcggcc	2280
gcagattata	aagatgacga	tgacaataa	taaggtacc			2319

&lt;210&gt; SEQ ID NO 160

&lt;211&gt; LENGTH: 1293

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 160

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gtcgacatga aaataaaaac aggtgcacgc atcctcgcat tatccgcatt aacgacgatg    60
atgttttccg cctcggtctc cgccaaaatc atcgaagccc gcacctcgga atccgacacg    120
gcagggccca acagcgacct ggacgtgaac actgacattt attccaaggt gctggtgact    180
gctatatacc tggcactctt cgtggtgggc actgtgggca actccgtgac agccttcaact    240
ctagcgcgga agaagtcact gcagagcctg cagagcactg tgcattacca cctgggcagc    300
ctggcaactg cggacctgct tatccttctg ctggccatgc cctgggagct atacaacttc    360
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cgtgatgcct gcacctatgc cacagccctc aatgtagcca gcctgagtgt ggagcgcctac    480
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caccagcgcc ccgagcaggg ccgagtgtgc accgtgggca cacacaacgg ttagagcac    840
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tacaacctgg tctccgcca cttccgccag gtctttctgt ccaogctggc ctgcctttgt   1140
cctgggtggc gccaccgccc aaagaagagg ccaacgttct ccaggaagcc caacagcatg   1200
tccagcaacc atgccttttc caccagcgcc acccgggaga ccctgtacgc ggccgcagat   1260
tataaagatg acgatgacaa ataataaggt acc                                     1293

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&lt;210&gt; SEQ ID NO 161

&lt;211&gt; LENGTH: 2652

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 161

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gtcgacatga aaataaaaac aggtgcacgc atcctcgcat tatccgcatt aacgacgatg    60
atgttttccg cctcggtctc cgccaaaatc gaagaaggta aactggtaat ctggattaac    120
ggcgataaag gctataacgg tctcgtgaa gtcggtaaga aattcgagaa agataaccgga    180
attaaagtoa ccgttgagca tccgataaaa ctggaagaga aattcccaca ggttgccgca    240
actggcgatg gccctgacat tatcttcttg gcacacgacc gctttggtgg ctacgtcaa    300
tctggcctgt tggctgaaat caccocggac aaagcgttcc aggacaagct gtatccgttt    360
accctgggatg ccgtacgtta caacggcaag ctgattgctt acccgatcgc tgttgaagcg    420
ttatcgctga tttataacaa agatctgctg ccgaaccgca caaaaacctg ggaagagatc    480
ccggcgctgg ataaagaact gaaagcgaaa ggtaagagcg cgctgatgtt caacctgcaa    540
gaacctgact tcacctggcc gctgattgct gctgacgggg gttatgcggt caagtatgaa    600
aacggcaagt acgacattaa agacgtgggc gtggataacg ctgggcgcaa agcgggtctg    660
accttctctg ttgacctgat taaaaacaaa cacatgaatg cagacaccga ttactocatc    720

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gcagaagctg cctttaataa aggcgaaaca gcgatgacca tcaacggccc gtgggcatgg 780
tccaacatcg acaccagcaa agtgaattat ggtgtaacgg tactgccgac cttcaagggt 840
caaccatcca aaccgttcgt tggcgtgctg agcgcaggta ttaacgccgc cagtccgaac 900
aaagagctgg cgaaagagtt cctcgaaaac tatctgctga ctgatgaagg tctggaagcg 960
gttaataaag acaaacccgt gggtgccgta gcgctgaagt cttacgagga agagttggcg 1020
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gacacggcag ggcccaacag cgacctggac gtgaacctg acatttattc caagtgctg 1200
gtgactgcta tatacctggc actctctgtg gtgggcaactc tgggcaactc cgtgacagcc 1260
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ggcagcctgg cactgtcggc cctgcttacc cttctgctgg ccatgccctg ggagctatac 1380
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gacggggcga tcctcgtcga tttctgggca gagtgggtgc gtcogtcaa aatgatcgcc 2400
ccgattctg atgaaatcgc tgacgaatat cagggcaaac tgaccgttg aaaaactgaa 2460
atcgatcaaa acctggcac tgcgcgaaa tatggcatcc gtggatccc gactctgctg 2520
ctgttcaaaa acggtgaagt ggcggcaacc aaagtgggtg cactgtctaa aggtcagttg 2580
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taataaggta cc 2652

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&lt;210&gt; SEQ ID NO 162

&lt;211&gt; LENGTH: 1626

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 162

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gtcgacatga aaataaaaac aggtgcacgc atcctcgcat tatccgcatt aacgacgatg      60
atgtttttccg cctcggctct cgcctaaatc atcgaagccc gcacctcgga atccgacacg      120
gcagggccca acagcgacct ggacgtgaac actgacattt attccaaggt gctggtgact      180
gctatatacc tggcactcct cgtggtgggc actgtgggca actccgtgac agccttcact      240
ctagcgcgga agaagtcaact gcagagcctg cagagcactg tgcattacca cctgggcagc      300
ctggcactgt cggacctgct tatccttctg ctggccatgc cgtggagct atacaacttc      360
atctgggtac accatccctg ggcccttggg gacgctggct gccgtggcta ctatttcctg      420
cgtgatgcct gcacctatgc cacagccctc aatgtagcca gcctgagtgt ggagcgctac      480
ttggccatct gccatccctt caaggccaag accctcatgt cccgcagccg caccaagaaa      540
ttcatcagtg ccatatggct agcttcggcg ctgctggcta taccatgct tttcaacctg      600
ggcctgcaga accgcagtgg tgacggcacg caccctggcg gcctggtgtg cacaccatt      660
gtggacacag ccactgtcaa ggtcgtcatc caggttaaca ccttcatgtc cttcctgttt      720
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ctgatgttct gctatatctc ggatgaacag tggactacgt tcctcttcga tttctaccac     1020
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cctgggtggc gccaccgcc aaagaagagg ccaacgttct ccaggaagcc caacagcatg     1200
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caaaacctg gcactgcgcc gaaatatggc atccgtggta tcccgaactct gctgctgttc     1500
aaaaacggtg aagtggcggc aaccaagtg ggtgcactgt ctaaaggta gttgaaagag     1560
ttcctcgacg ctaacctggc agcggccgca gattataaag atgacgatga caaataataa     1620
ggtacc                                             1626

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&lt;210&gt; SEQ ID NO 163

&lt;211&gt; LENGTH: 26

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Cloning primer

&lt;400&gt; SEQUENCE: 163

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ggtgcacgca tcctcgcat atccgc                                             26

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&lt;210&gt; SEQ ID NO 164

&lt;211&gt; LENGTH: 31

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Cloning primer



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<400> SEQUENCE: 164  
cgcacggcat accagaaagc ggacatctgc g 31

<210> SEQ ID NO 165  
<211> LENGTH: 43  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 165  
ccgcggtcga catgaaaata aaaacaggtg cacgcatcct cgc 43

<210> SEQ ID NO 166  
<211> LENGTH: 66  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 166  
gccgtgtcgg attccgaggt gcggccttcg atacgcacgg cataccaaga aagcgggatg 60  
ttcggc 66

<210> SEQ ID NO 167  
<211> LENGTH: 24  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 167  
cctcggaatc cgacacggca gggc 24

<210> SEQ ID NO 168  
<211> LENGTH: 26  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 168  
gtacagggtc tcccgggtgg cgctgg 26

<210> SEQ ID NO 169  
<211> LENGTH: 43  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 169  
ccgcgatcga aggccgcacc tcggaatccg acacggcagg gcc 43

<210> SEQ ID NO 170  
<211> LENGTH: 73  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 170

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ggcgcggtac ctttgcctc gtcctcttta taatctgcgg ccgctacag ggtctccgg 60  
gtggcgctgg tgg 73

<210> SEQ ID NO 171  
<211> LENGTH: 51  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 171

ggcgcggtac cttattattt gtcctcgtca tttttataat ctggggccgc g 51

<210> SEQ ID NO 172  
<211> LENGTH: 79  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 172

ccgcattaac gacgatgatg ttttccgct cggtctcgc caaatcctc gaaggccga 60  
cctcggaatc cgacacggc 79

<210> SEQ ID NO 173  
<211> LENGTH: 82  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 173

ccgcggtcga catgaaaata aaaacagtg cagcctcct cgcattatcc gcattaacga 60  
cgatgatggt ttccgctcgc gc 82

<210> SEQ ID NO 174  
<211> LENGTH: 33  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 174

ccgcgagcga taaaattatt cacctgactg acg 33

<210> SEQ ID NO 175  
<211> LENGTH: 39  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 175

gcccgccagg ttagcgtcga ggaactcttt caactgacc 39

<210> SEQ ID NO 176  
<211> LENGTH: 37  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

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&lt;400&gt; SEQUENCE: 176

gcggccgcaa gcgataaaat tattcacctg actgacg 37

&lt;210&gt; SEQ ID NO 177

&lt;211&gt; LENGTH: 38

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Cloning primer

&lt;400&gt; SEQUENCE: 177

ggcgctgagg ccgcatcatc atgatcttta taatcgcc 38

&lt;210&gt; SEQ ID NO 178

&lt;211&gt; LENGTH: 2465

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 178

gtcgacatgg ggcaaccggg gaacggcagc gccttcttgc tggcaccxaa tggagccat 60

gcgcccggacc acgacgtcac gcagcaaagg gacgaggtgt ggggtgtggg catgggcatc 120

gtcatgtctc tcatcgtcct ggccatcgtg tttggcaatg tgetgtcat cacagccatt 180

gccaaagtgg agcgtctgca gacggtcacc aactacttca tcaactcact gcctgtgct 240

gatctgtgca tgggcctagc agtggtgccc tttggggccg cccatattct tatgaaaatg 300

tggacttttg gcaacttctg gtgagagttt tggacttcca ttgatgtgct gtgctgcaag 360

gccagcattg agaccctgtg cgtgatcgca gtggatcgtc actttgcat tacttcacct 420

ttcaagtacc agagcctgct gaccaagaat aaggcccggg tgatcattct gatggtgtgg 480

attgtgtcag gccttayctc cttcttgccc attcagatgc actggtacag ggccaccac 540

caggaaagca tcaactgcta tgccaatgag acctgctgtg acttcttcac gaaccaagcc 600

tatgccattg cctcttccat cgtgtccttc tacgttcccc tggatgatcat ggtcttcgtc 660

tactccaggg tctttcagga ggccaaaagg cagctccaga agattgacaa atctgagggc 720

cgcttccatg tccagaacct tagccagggt gagcaggatg ggcggacggg gcatggactc 780

cgcagatctt ccaagttctg cttgaaggag cacaagccc tcaagcgtt aggcacatc 840

atgggcactt tcaccctctg ctggctgccc ttcttcatcg ttaacattgt gcatgtgatc 900

caggataacc tcatccgtaa ggaagtttac atcctcctaa attggatagg ctatgtcaat 960

tctgtttcoa atccccttat ctactgccgg agcccagatt tcaggattgc cttccaggag 1020

cttctgtgoc tgcgcaggtc ttctttgaag gcctatggca atggctactc cagcaacggc 1080

aacacagggg agcagagtgg atatcacgtg gaacaggaga aagaaaataa actgctgtgt 1140

gaagacctcc caggcacgga agactttgtg ggccatcaag gtactgtgoc tagcgataac 1200

attgattcac aaggaggaa ttgtagtaca aatgactcac tgctagagcg tggccagacg 1260

gtcaccaccc tgcagctcga gggctgcctc gggaaacagta agaccgagga ccagcgcaac 1320

gaggagaagg cgcagcgtga ggccaacaaa aagatcgaga agcagctgca gaaggacaag 1380

caggtctaac gggccacgca ccgctgctg ctgctgggtg ctggagaatc tggtaaaagc 1440

accattgtga agcagatgag gatcctgcat gtaaatgggt ttaatggaga cagtgagaag 1500

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gcaaccaaag tgcaggacat caaaaacaac ctgaaagagg cgattgaaac cattgtggcc 1560
gccatgagca acctggtgcc ccccgtaggag ctggccaacc cggagaacca gttcagagtg 1620
gactacatcc tgagtgtgat gaacgtgcct gactttgact tccctcccga attctatgag 1680
catgccaaag ctctgtggga ggatgaagga gtgcgtgcct gctacgaacg ctccaacgag 1740
taccagctga ttgactgtgc ccagtacttc ctggacaaga tgcagctgat caagcaggct 1800
gactatgtgc cgagcgatca ggacctgctt cgctgccgtg tcctgacttc tggaaatcttt 1860
gagaccaagt tccaggtgga caaagtcaac ttccacatgt ttgacgtggg tggccagcgc 1920
gatgaacgcc gcaagtggat ccagtgtctc aacgatgtga ctgccatcat cttcgtgggtg 1980
gccagcagca gctacaacat ggtcatccgg gaggacaacc agaccaaccg cctgcaggag 2040
gctctgaacc tcttcaagag catctggaac aacagatggc tgcgcacccat ctctgtgatc 2100
ctgttcctca acaagcaaga tctgctcgct gagaaagtcc ttgctgggaa atcgaagatt 2160
gaggactact ttccagaatt tgctcgtac actactcctg aggatgctac tcccgagccc 2220
ggagaggacc cacgcgtgac cggggccaag tacttcattc gagatgagtt tctgaggatc 2280
agcactgcca gtggagatgg gcgtcactac tgctaccctc atttcaactg cgctgtggac 2340
actgagaaca tccgccgtgt gttcaacgac tgccgtgaca tcattcagcg catgcacctt 2400
cgtcagtacg agctgctcat cgattaataa tctagaggat ccccgcgccc tcatccgaaa 2460
gggcg 2465

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&lt;210&gt; SEQ ID NO 179

&lt;211&gt; LENGTH: 2485

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 179

```

gtcgacatgg ggcaaccctg gaacgcagc gccttcttgc tggcaccocaa tggaaagccat 60
gcgcccggacc acgacgtcac gcagcaaaagg gacgaggtgt ggggtgtggg catgggcatc 120
gtcatgtctc tcatcgtcct ggccatcgtg tttggcaatg tgctggtcat cacagccatt 180
gccaaagtctg agcgtctgca gacggtcacc aactacttca tcacttcaact ggcctgtgct 240
gatctgtgca tggccctagc agtggtgccc tttggggccg cccatattct tatgaaaatg 300
tggacttttg gcaacttctg gtgcgagttt tggacttcca ttgatgtgct gtgcgtcacg 360
gccagcattg agaccctgtg cgtgatcgca gtggatcgtc actttgccat tacttcaact 420
ttcaagtacc agagcctgct gaccaagaat aaggcccggg tgatcattct gatggtgtgg 480
attgtgtcag gccttayctc cttcttggcc attcagatgc actggtacag ggccaaccac 540
caggaagcca tcaactgcta tgccaatgag acctgctgtg acttcttcac gaaccaagcc 600
tatgccattg cctcttccat cgtgtccttc tacgttcccc tggatgatcat ggtcttcgtc 660
tactccaggg tctttcagga ggccaaaagg cagctccaga agattgacaa atctgagggc 720
cgcttccatg tccagaacct tagccaggtg gagcaggatg ggcggacggg gcatggactc 780
cgcagatctt ccaagttctg cttgaaggag cacaaagccc tcaagacgtt aggcacatc 840
atgggcactt tcacctctg ctggctgccc ttcttcatcg ttaacattgt gcatgtgatc 900
caggataacc tcatccgtaa ggaagtttac atcctcctaa attggatag ctatgtcaat 960

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tctggttca atcccccttat ctactgccgg agcccagatt tcaggattgc cttccaggag 1020
cttctgtgcc tgcgcaggtc ttctttgaag gcctatggca atggctactc cagcaacggc 1080
aacacagggg agcagagtgg atatcacgtg gaacaggaga aagaaaataa actgctgtgt 1140
gaagacctcc caggcacgga agactttgtg ggccatcaag gtactgtgcc tagcgataac 1200
attgattcac aaggaggaa ttgtagtaca aatgactcac tgctagagcg tggccagacg 1260
gtcaccaacc tgcagtaata atcaaggagg ccctcgagat gggctgctc gggaacagta 1320
agaccgagga ccagcgcaac gaggagaagg cgcagcgtga ggccaacaaa aagatcgaga 1380
agcagctgca gaagacaag caggtctacc gggccacgca ccgctgctg ctgctgggtg 1440
ctggagaatc tggtaaaagc accattgtga agcagatgag gatcctgcat gttaatgggt 1500
ttaatggaga cagtgagaag gcaacaaaag tgcaggacat caaaaaaac ctgaaagagg 1560
cgattgaaac cattgtggcc gccatgagca acctggtgcc ccccgaggag ctggccaacc 1620
ccgagaacca gttcagagtg gactacatcc tgagtgtgat gaacgtgcct gactttgact 1680
tcccctccga attctatgag catgccaaag ctctgtggga ggatgaagga gtgcgtgcct 1740
gctacgaaoc ctccaacgag taccagctga ttgactgtgc ccagtacttc ctggacaaga 1800
tcgacgtgat caagcaggct gactatgtgc cgagcgatca ggacctgctt cgctgccgtg 1860
tctctgacttc tggaaatcttt gagaccaagt tccaggtgga caaagtcaac ttccacatgt 1920
ttgacgtggg tggccagcgc gatgaacgcc gcaagtggat ccagtgcttc aacgatgtga 1980
ctgccatcat ctctgtggtg gccagcagca gctacaacat ggtcatcccg gaggacaacc 2040
agaccaaccg cctgcaggag gctctgaacc tcttcaagag catctggaac aacagatggc 2100
tgccaccatc ctctgtgatc ctgttctca acaagcaaga tctgctcgct gagaaagtcc 2160
ttgctgtggaa atcgaagatt gaggactact ttccagaatt tgctcgctac actactcctg 2220
aggatgctac tcccagagccc ggagaggacc cacgcgtgac ccgggccaag tacttcattc 2280
gagatgagtt tctgaggatc agcaactgcca gtggagatgg gcgtcactac tgctaccctc 2340
atctacactg cgctgtggac actgagaaca tccgccgtgt gttcaacgac tgccgtgaca 2400
tcattcagcg catgcacctt cgtcagtagc agctgctcat cgattaataa tctagaggat 2460
ccccgcgccc tcatccgaaa gggcg 2485

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&lt;210&gt; SEQ ID NO 180

&lt;211&gt; LENGTH: 1146

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Homo sapien

&lt;400&gt; SEQUENCE: 180

```

ctcgagatgg gctgcctcgg gaacagtaag accgaggacc agcgaacga ggagaaggcg 60
cagcgtgagg ccaacaaaaa gatcgagaag cagctgcaga aggacaagca ggtctaccgg 120
gccacgcacc gcctgctgct gctgggtgct ggagaatctg gtaaaagcac cattgtgaag 180
cagatgagga tcctgcatgt taatgggttt aatggagaca gtgagaaggc aaccaaagtg 240
caggacatca aaaacaacct gaaagaggcg attgaaacca ttgtggccgc catgagcaac 300
ctggtgcccc ccgtggagct ggccaacccc gagaaccagt tcagagtgga ctacatcctg 360
agtgtgatga acgtgcctga ctttgacttc cctcccgaat tctatgagca tgccaaggct 420
ctgtggggagg atgaaggagt gcgtgcctgc tacgaacgct ccaacgagta ccagctgatt 480

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gactgtgccc agtacttccct ggacaagatc gacgtgatca agcaggctga ctatgtgccg 540
agcgatcagg acctgcttcg ctgccgtgtc ctgacttctg gaatctttga gaccaagttc 600
caggtggaca aagtcaactt ccacatgttt gacgtgggtg gccagcgcga tgaacgccgc 660
aagtggatcc agtgcttcaa cgatgtgact gccatcatct tcgtggggc cagcagcagc 720
tacaacatgg tcatccggga ggacaaccag accaaccgcc tgcaggaggc tctgaacctc 780
ttcaagagca tctggaacaa cagatggctg cgcaccatct ctgtgatcct gttcctcaac 840
aagcaagatc tgctcgtga gaaagtcctt gctgggaaat cgaagattga ggactacttt 900
ccagaatttg ctgctacac tactcctgag gatgctactc ccgagcccg agaggacca 960
cgctgaccc gggccaagta cttcattcga gatgagtttc tgaggatcag cactgccagt 1020
ggagatggg gtcactactg ctaccctcat ttcacctgcg ctgtggacac tgagaacatc 1080
cgccgtgtgt tcaacgactg ccgtgacatc attcagcga tgcaccttcg tcagtacgag 1140
ctgctc 1146

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<210> SEQ ID NO 181
<211> LENGTH: 1194
<212> TYPE: DNA
<213> ORGANISM: Homo sapien

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<400> SEQUENCE: 181

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ctcgagatgg gctgcctcgg gaacagtaag accgaggacc agcgcaacga ggagaaggcg 60
cagcgtgagg ccaacaaaaa gatcgagaag cagctgcaga aggacaagca ggtctaccgg 120
gccacgcacc gcctgctgct gctgggtgct ggagaatctg gtaaaagcac cattgtgaag 180
cagatgagga tcctgcatgt taatgggttt aatggagagg gcggcgaaga ggaccgcag 240
gctgcaagga gcaacagcga tggtgagaag gcaaccaaag tgcaggacat caaaaacaac 300
ctgaaagagg cgattgaaac cattgtggcc gccatgagca acctggtgcc ccccggtggag 360
ctggccaacc ccgagaacca gttcagatg gactacatcc tgagtgtgat gaacgtgcct 420
gactttgact tccctcccga attctatgag catgccaaag ctctgtggga ggatgaagga 480
gtgcgtgcct gtacgaacg ctccaacgag taccagctga ttgactgtgc ccagtacttc 540
ctggacaaga tcgacgtgat caagcaggct gactatgtgc cgagcgatca ggacctgctt 600
cgctgccgtg tcctgacttc tggaatcttt gagaccaagt tccaggtgga caaagtcaac 660
ttccacatgt ttgacgtggg tggccagcgc gatgaacgcc gcaagtggat ccagtgcttc 720
aacgatgtga ctgccatcat ctctgtggtg gccagcagca gctacaacat ggtcatccgg 780
gaggacaacc agaccaaccg cctgcaggag gctctgaacc tcttcaagag catctggaac 840
aacagatggc tgcgcacat ctctgtgato ctgttcctca acaagcaaga tctgctcgt 900
gagaaagtcc ttgtgggaa atcgaagatt gaggactact ttccagaatt tgctcgtac 960
actactcctg aggatgctac tcccagccc ggagaggacc cacgcgtgac ccgggccaag 1020
tacttcattc gagatgagtt tctgaggatc agcactgcca gtggagatgg gcgtcactac 1080
tgctaccctc atttcacctg cgctgtggac actgagaaca tccgccgtgt gttcaacgac 1140
tgccgtgaca tcattcagcg catgcacctt cgtcagtacg agctgctcat cgat 1194

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<210> SEQ ID NO 182
<211> LENGTH: 1089
<212> TYPE: DNA
<213> ORGANISM: Homo sapien

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&lt;400&gt; SEQUENCE: 182

ctcgagatga ctctggagtc catcatggcg tgctgcctga gcgaggagcc caaggaagcc	60
cggcggatca acgacgagat cgagcggcag ctccgcaggg acaagcggga cgcccgccgg	120
gagctcaagc tgctgctgct cgggacagga gagagtggca agagtacgtt tatcaagcag	180
atgagaatca tccatgggctc aggatactct gatgaagata aaaggggctt caccaagctg	240
gtgtatcaga acatcttcac ggccatgcag gccatgatca gagccatgga cacactcaag	300
atcccataca agtatgagca caataaggct catgcacaat tagttcgaga agttgatgtg	360
gagaagggtg ctgcttttga gaatccatat gtagatgcaa taaagagttt atggaatgat	420
cctggaatcc aggaatgcta tgatagacga cgagaatata aattatctga ctctacaaa	480
tactatctta atgacttggc ccgctgagct gacctgcct acctgcctac gcaacaagat	540
gtgcttagag ttcagagccc caccacaggg atcatcgaat acctcttga cttacaaagt	600
gtcattttca gaatggtcga ttagggggc caaaggcag agagaagaaa atggatacac	660
tgctttgaaa atgtcacctc tatcatgttt ctagtgcgc ttagtgaata tgatcaagtt	720
ctcgtggagt cagacaatga gaaccgaatg gagaaagca aggtctctt tagaacaatt	780
atcacatacc cctgggtcca gaactcctcg gttattctgt tcttaacaa gaaagatctt	840
ctagaggaga aaatcatgta ttcccatcta gtcgactact tcccagaata tgatggacc	900
cagagagatg cccagcgagc ccgagaatcc attctgaaga tgttcgtgga cctgaacca	960
gacagtgaca aaattatcta ctcccacttc acgtgcgcca cagacaccga gaatatccgc	1020
tttgccttg ctgcccgtcaa ggacaccatc ctccagttga acctgaagga gtacaatctg	1080
gtcatcgat	1089

&lt;210&gt; SEQ ID NO 183

&lt;211&gt; LENGTH: 1077

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Homo sapien

&lt;400&gt; SEQUENCE: 183

ctcgagatgg gctgcaccgt gagcggcag gacaaggcgg cggccgagcg ctctaagatg	60
atcgacaaga acctgcggga ggacggagag aaggcggcgc gggagtgaa gttgctgctg	120
ttgggtgctg gggagttagg gaagagcacc atcgtcaagc agatgaagat catccacgag	180
gatggctact ccgaggagga atgccggcag taccgggagg ttgtctacag caacaccatc	240
cagtccatca tggccattgt caaagccatg ggaaacctgc agatcgactt tgccgacccc	300
tccagagcgg acgacgccag gcagctattt gcaactgtcct gcaccgccga ggagcaaggc	360
gtgctccctg atgacctgtc cggcgtcatc cggaggctct gggctgacca tgggtgtagc	420
gcctgcttgg gccgctcaag ggaataccag ctcaacgact cagctgccta ctacctgaac	480
gacctggagc gtattgcaca gagtgactac atccccacac agcaagatgt gctacggacc	540
cgcgtaaaga ccacggggat cgtggagaca cacttcacct tcaaggacct acacttcaag	600
atgtttgatg tgggtgtgca gcggtctgag cggaagaagt ggtaccactg ctttgagggc	660
gtcacagcoa tcatcttctg cgtagccttg agcgcctatg acttgggtgct agctgaggac	720
gaggagatga accgcatgca tgagagcatg aagctattcg atagcatctg caacaacaag	780
tggttcacag acacgtccat catcctcttc ctcaacaaga aggacctgtt tgaggagaag	840

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atcacacaca gtcccctgac catctgcttc cctgagtaca caggggccaa caaatatgat    900
gaggcagcca gctacatcca gagtaagttt gaggacctga ataagcgcaa agacaccaag    960
gagatctaca cgcacttcac gtgcccacc gacaccaaga acgtgcagtt cgtgtttgac    1020
gccgtcaccg atgtcatcat caagaacaac ctgaaggact gcggcctctt catgcat      1077

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<210> SEQ ID NO 184

<211> LENGTH: 1155

<212> TYPE: DNA

<213> ORGANISM: Homo sapien

<400> SEQUENCE: 184

```

ctcgagatgt ccggggtggt gcggaccctc agccgctgcc tgctgccggc cgaggccggc    60
ggggcccgcg agcgcagggc gggcagcggc gcgcgcgacg cggagcgcga ggcccggagg    120
cgtagccgcg acatcgacgc gctgctggcc cgcgagcggc gcgcggtccg gcgcctggtg    180
aagatcctcg tgctggggcg gggcgagagc ggcaagtcca cgttctctca gcagatgcgc    240
atcatccacg gccgcgagtt cgaccagaag gcgctgctgg agttccgcga caccatcttc    300
gacaacatcc tcaagggctc aagggttctt gttgatgcac gagataagct tggcattcct    360
tggcagtatt ctgaaaatga gaagcatggg atgttctctga tggccttcga gaacaaggcg    420
gggctgcctg tggagccggc caccttccag ctgtacgtcc cggccctgag cgcactctgg    480
agggattctg gcatcagggg ggctttcagc cggagaagcg agtttcagct gggggagtcg    540
gtgaagtact tcctggacaa ctgggaccgg atcggccagc tgaattactt tcctagtaag    600
caagatatcc tgctggctag gaaagccacc aagggaaattg tggagcatga cttcgttatt    660
aagaagatcc cctttaagat ggtggatggt ggcggccagc ggtcccagcg ccagaagtgg    720
ttccagtgtg tcgacgggat cacgtccatc ctgttcatgg tctctccag cgagtacgac    780
caggtcctca tggagcagac gcgcaccaac cggctggtgg agtccatgaa catcttcgag    840
accatcgtea acaacaagct cttcttcaac gtctccatca ttctcttctt caacaagatg    900
gacctccttg tggagaaggt gaagaccgtg agcatcaaga agcacttccc ggacttcagg    960
ggcgaccocg accagctgga ggacgtccag cgctacctgg tccagtgtct cgacaggaag   1020
agacggaacc gcagcaagcc actcttccac cacttcacca ccgccatcga caccgagaac   1080
gtccgcttgc tgttccatgc tgtgaaagac accatcctgc aggagaacct gaaggacatc   1140
atgctgcaga tcgat                                     1155

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<210> SEQ ID NO 185

<211> LENGTH: 3307

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 185

```

gtcgacatgg ggcaaccggg gaacggcagc gccttcttgc tggcaccxaa tgggaagccat    60
gcgcccggacc acgacgtcac gcagcaaagg gacgaggtgt ggggtggtggg catgggcatc    120
gtcatgtctc tcacgtctct ggccatcgtg tttggcaatg tgctggtcat cacagccatt    180
gccaaagtto agcgtctgca gaocgtcacc aactacttca tcaottcact ggctgtgct    240
gatctgtgta tgggcctagc agtggtgccc tttggggccg cccatattct tatgaaaatg    300
tggacttttg gcaacttctg gtgcgagttt tggacttcca ttgatgtgct gtgcgtcacg    360

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gccagcattg agaccctgtg cgtgatcgca gtggatcgct actttgccat tacttcacct	420
ttcaagtacc agagcctgct gaccaagaat aaggcccggg tgatcattct gatggtgtgg	480
attgtgtcag gccttayctc cttcttgccc attcagatgc actggtacag gcccaccac	540
caggaagcca tcaactgcta tgccaatgag acctgctgtg acttcttcac gaaccaagcc	600
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&lt;210&gt; SEQ ID NO 186

&lt;211&gt; LENGTH: 3284

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 186

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&lt;210&gt; SEQ ID NO 187

&lt;211&gt; LENGTH: 25

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Cloning primer

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<400> SEQUENCE: 187

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<211> LENGTH: 30

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 188

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<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 189

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<210> SEQ ID NO 190

<211> LENGTH: 67

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 190

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<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 191

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<211> LENGTH: 29

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 192

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<211> LENGTH: 44

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 193

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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 194

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<210> SEQ ID NO 195  
<211> LENGTH: 37  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 195

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<210> SEQ ID NO 196  
<211> LENGTH: 26  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 196

ccgggggtggt gcggaccctc agccgc 26

<210> SEQ ID NO 197  
<211> LENGTH: 28  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 197

ctgcagcatg atgtccttca ggttctcc 28

<210> SEQ ID NO 198  
<211> LENGTH: 41  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 198

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<210> SEQ ID NO 199  
<211> LENGTH: 39  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 199

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<210> SEQ ID NO 200

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<211> LENGTH: 28  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
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<210> SEQ ID NO 201  
<211> LENGTH: 29  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 201  
  
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<211> LENGTH: 30  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 202  
  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
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<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 204  
  
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<210> SEQ ID NO 205  
<211> LENGTH: 30  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer  
  
<400> SEQUENCE: 205  
  
ccttcagggtt gttcttgatg atgacatcgg 30  
  
<210> SEQ ID NO 206  
<211> LENGTH: 31  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:

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<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 206

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<210> SEQ ID NO 207  
<211> LENGTH: 55  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 207

gcgccatcga tgaagaggcc gcagtccttc aggttggttct tgatgatgac atcgg 55

<210> SEQ ID NO 208  
<211> LENGTH: 29  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 208

gggctgcctc gggaacagta agaccgag 29

<210> SEQ ID NO 209  
<211> LENGTH: 29  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 209

gagcagctcg tactgacgaa ggtgcatgc 29

<210> SEQ ID NO 210  
<211> LENGTH: 31  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 210

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<211> LENGTH: 40  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 211

gcgccatcga tgagcagctc gtactgacga aggtgcatgc 40

<210> SEQ ID NO 212  
<211> LENGTH: 107  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 212

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ggctcgaggg cctccttgat tattactcga gggcctcctt gattattact gcaggttggt 60

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<210> SEQ ID NO 213

<211> LENGTH: 61

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 213

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c 61

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<211> LENGTH: 38

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 214

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<210> SEQ ID NO 215

<211> LENGTH: 29

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 215

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<210> SEQ ID NO 216

<211> LENGTH: 52

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 216

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<210> SEQ ID NO 217

<211> LENGTH: 51

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 217

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<210> SEQ ID NO 218

<211> LENGTH: 32

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 218



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ggctgtgggt agaagtgaaa cggggtttac cg 32

<210> SEQ ID NO 219  
<211> LENGTH: 32  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 219

ctttaccata taatgctccc ttgtttaa ac ag 32

<210> SEQ ID NO 220  
<211> LENGTH: 43  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 220

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<210> SEQ ID NO 221  
<211> LENGTH: 60  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 221

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<210> SEQ ID NO 222  
<211> LENGTH: 27  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 222

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<210> SEQ ID NO 223  
<211> LENGTH: 27  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
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<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 223

ccagaccaac tgtaatggt agcgacc 27

<210> SEQ ID NO 224  
<211> LENGTH: 35  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 224

ggtaaagacc atgattacgg attcactggc cgtcg 35

<210> SEQ ID NO 225

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<211> LENGTH: 77  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 225

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<210> SEQ ID NO 226  
<211> LENGTH: 89  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 226

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cgatgatggt ttccgcctcg gctctcgcc 89

<210> SEQ ID NO 227  
<211> LENGTH: 69  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 227

cgctgaccga ggctgcagg cgggcttcga tgattttggc gagagccgag gcggaaaaca 60  
tcctcgtcg 69

<210> SEQ ID NO 228  
<211> LENGTH: 81  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 228

cgaagcccgc ctgcaggcct cggctcagc cgaatctaga gattataaag atgacgatga 60  
caaataataa gctagcggcg c 81

<210> SEQ ID NO 229  
<211> LENGTH: 27  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 229

gcgccgctag cttattatgt gtcctcg 27

<210> SEQ ID NO 230  
<211> LENGTH: 26  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 230

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ggtgcacgca tcctcgatt atccg 26

<210> SEQ ID NO 231  
<211> LENGTH: 30  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 231

ggcgttttcc atggtggcgg caatacgtgg 30

<210> SEQ ID NO 232  
<211> LENGTH: 52  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 232

cgcggtatgca tatgaaaata aaaacaggtg cacgcaccc cgcattatcc gc 52

<210> SEQ ID NO 233  
<211> LENGTH: 82  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 233

ccgagcctg caggcggcct tcgatacgca cggcatacca gaaagcggac tggcggttt 60

ccatggtggc ggcaatacgt gg 82

<210> SEQ ID NO 234  
<211> LENGTH: 87  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 234

gcgccgtag cttattatgt gtcacgtca tctttataat ctctagattc ggcgtcgacc 60

gaggcctgca ggcgggcttc gatacgc 87

<210> SEQ ID NO 235  
<211> LENGTH: 26  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 235

cctgactgac gacagttttg acacgg 26

<210> SEQ ID NO 236  
<211> LENGTH: 32  
<212> TYPE: DNA  
<213> ORGANISM: Artificial Sequence  
<220> FEATURE:  
<223> OTHER INFORMATION: Cloning primer

<400> SEQUENCE: 236

-continued

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cctttagaca gtgcacccac ttgtgtgccc gc 32

<210> SEQ ID NO 237  
 <211> LENGTH: 78  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

&lt;400&gt; SEQUENCE: 237

cgcggtgaca ggcctcggtc gacgccgaat ctagaagcga taaaattatt cacctgactg 60

acgacagttt tgacacgg 78

<210> SEQ ID NO 238  
 <211> LENGTH: 95  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Cloning primer

&lt;400&gt; SEQUENCE: 238

gcgccgctag cttattatgt gtcacgtca tctttataat ccgccagggt ctctttcaac 60

tgacctttag acagtgcacc cactttggtt gccgc 95

<210> SEQ ID NO 239  
 <211> LENGTH: 216  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 239

gaattcaggc gctttttaga ctggtcgtaa tgaaattcag gaggttctgc atatgaaaat 60

aaaaacaggt gcacgcattc tcgattatc cgcattaacg acgatgatgt ttccgcctc 120

ggctctcgcc aaaatcatcg aagcccgctc gcaggcctcg gtcgacgccc aatctagaga 180

ttataaagat gacgatgaca aataataagc tagagg 216

<210> SEQ ID NO 240  
 <211> LENGTH: 202  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 240

ccatacccgt ttttttgggc tagcaggagg ccctgcatat gaaaataaaa acaggtgcac 60

gcatcctcgc attatccgca ttaacgacga tgatgttttc cgctcggct ctcgccaaaa 120

tcatcgaagc ccgctcgac gcctcggtcg acgccgaatc tagagattat aaagatgacg 180

atgacaaaata ataagctaga gg 202

<210> SEQ ID NO 241  
 <211> LENGTH: 182  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 241

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```

aggaggttct gcatatgaaa ataaaaacag gtgcacgcat cctcgatta tccgcattaa 60
cgacgatgat gttttccgcc tcggctctcg ccaaatcat cgaagccgc ctgcaggcct 120
cggtcgacgc cgaatctaga gattataaag atgacgatga caaataataa gctagaggta 180
cc 182

```

```

<210> SEQ ID NO 242
<211> LENGTH: 182
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Fusion vector

```

```
<400> SEQUENCE: 242
```

```

aggaggttct gcatatgaaa ataaaaacag gtgcacgcat cctcgatta tccgcattaa 60
cgacgatgat gttttccgcc tcggctctcg ccaaatcat cgaagccgc ctgcaggcct 120
cggtcgacgc cgaatctaga gattataaag atgacgatga caaataataa gctagaggta 180
cc 182

```

```

<210> SEQ ID NO 243
<211> LENGTH: 1080
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Fusion vector

```

```
<400> SEQUENCE: 243
```

```

gaattcagcg gctttttaga ctggctgtaa tgaaattcag gaggttctgc atatgaaat 60
aaaaacaggt gcacgcattc tcgcattatc cgcattaacg acgatgatgt tttccgcctc 120
ggctctcgcc aaaatcgaag aaggtaaact ggtaactctgg attaacggcg ataaaggcta 180
taacggctct gctgaagtcg gtaagaaatt cgagaaagat accggaatta aagtcaccgt 240
tgagcatccg gataaaactg aagagaaatt cccacaggtt gcggaactg gcgatggccc 300
tgacattatc ttctgggac acgaccgctt tgggtgctac gctcaatctg gctgttggc 360
tgaaatcacc ccggacaag cgttccagga caagctgtat ccgtttacct gggatgccc 420
acgttaacaac ggcaagctga ttgcttacc gatcgtgtt gaagcgttat cgctgattta 480
taacaaagat ctgctgccga accgcgcaaa aacctgggaa gagatcccgg cgctggataa 540
agaactgaaa gcgaaaggta agagcgcgct gatgttcaac ctgcaagaac cgtacttcac 600
ctggccgctg attgctgctg acggggggtta tgcgttcaag tatgaaaacg gcaagtacga 660
cattaaagac gtggcgctgg ataacgctgg cgcgaaagcg ggtctgacct tcctggttga 720
cctgattaaa aacaacaca tgaatgcaga caccgattac tccatcgag aagctgcctt 780
taataaaggc gaaacagcga tgaccatcaa cggcccgtgg gcatggcca acatcgacac 840
cagcaaagtg aattatggtg taacggtact gccgacctc aaggtcaac catccaaacc 900
gttcgttggc gtgctgagcg caggtattaa cgccgccagt ccgaacaaag agctggcgaa 960
agagttcctc gaaaactatc tgctgactga tgaaggtctg gaagcgggta ataaagacaa 1020
accgctgggt gccgtagcgc tgaagtctta cgaggaagag ttggcgaaa atccaogtat 1080

```

```

<210> SEQ ID NO 244
<211> LENGTH: 1196
<212> TYPE: DNA

```

-continued

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 244

```

ccataccggt ttttttggc tagcaggagg cctgcatat gaaaataaaa acaggtgcac    60
gcatcctcgc attatccgca ttaacgacga tgatgttttc cgctcggct ctgccaaaa    120
tcgaagaagg taaactggta atctggatta acggcgataa aggctataac ggtctcgtg    180
aagtcggtaa gaaattcag aaagataccg gaattaaagt caccgttgag catccggata    240
aactggaaga gaaattccca caggttgccg caactggcga tggccctgac attatcttct    300
gggcacacga ccgctttggt ggctacgctc aatctggcct gttggctgaa atcaccocgg    360
acaaagcgtt ccaggacaag ctgtatccgt ttacctggga tgcogtacgt tacaacggca    420
agctgattgc ttaccgcatc gctgttgaag cgttatcgt gatttataac aaagatctgc    480
tgccgaaccc gccaaaaacc tgggaagaga tcccggcgct ggataaagaa ctgaaagcga    540
aaggtaaagc cgcgctgatg ttcaacctgc aagaaccgta cttcacctgg ccgctgattg    600
ctgctgacgg gggttatgcy ttcaagtatg aaaacggcaa gtacgacatt aaagacgtgg    660
gcgtggataa cgctggcgcg aaagcgggtc tgaccttctt gttgacctg attaaaaaca    720
aacacatgaa tgcagacacc gattactcca tcgcagaagc tgcctttaat aaaggcga    780
cagcgtgatc catcaacggc ccgtgggcat ggtccaacat cgacaccagc aaagtgaatt    840
atggtgtaac ggtactgccg acctcaagg gtcaaccatc caaacggtt gttggcgtgc    900
tgagcgcagg tattaacgcc gccagtccga acaaagagct ggcgaaagag ttcctcgaaa    960
actatctgct gactgatgaa ggtctggaag cggtaataa agacaaaccg ctgggtgccg   1020
tagcgcgtgaa gtcttacgag gaagagttgg cgaaagatcc acgtattgcc gccaccatgg   1080
aaaacgcccc gtccgctttc tggatgccc tgcgtatcga agcccgcctg caggcctcgg   1140
tcgacgcccga atctagagat tataaagatg acgatgacaa ataataagct agagga     1196

```

&lt;210&gt; SEQ ID NO 245

&lt;211&gt; LENGTH: 1171

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 245

```

aggaggttct gcatatgaaa ataaaaacag gtgcacgcat cctcgatta tccgattaa    60
cgacgatgat gttttccgcc tcggctctcg ccaaaatcga agaaggtaaa ctggtaatct   120
ggattaacgg cgataaaggc tataacggtc tcgctgaagt cggtaagaaa ttcgagaaag   180
ataccggaat taaagtccac gttgagcatc cggataaact ggaagagaaa tccccacagg   240
ttgcggcaac tggcgtggc cctgacatta tttctgggc acacgaccgc tttggtggct   300
acgctcaatc tggcctggtg gctgaaatca ccccgacaa agcgttccag gacaagctgt   360
atccgtttac ctgggatgcc gtacgttaca acggcaagct gattgcttac ccgatcgtg   420
ttgaagcgtt atcgtgatt tataacaaag atctgctgcc gaaccgccca aaaacctggg   480
aagagatccc ggcgctggat aaagaactga aagcgaaagg taagagcgcg ctgatgttca   540
acctgcaaga accgtacttc acctggccgc tgattgctgc tgacgggggt tatgcgttca   600

```

## -continued

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```

agtatgaaaa cggcaagtac gacattaaag acgtgggctt ggataacgct ggcgcgaaaag 660
cgggtctgac cttcctgggt gacctgatta aaaacaaaca catgaatgca gacaccgatt 720
actccatcgc agaagctgcc ttaataaaag gcgaaacagc gatgaccatc aacggcccgt 780
gggcatggtc caacatcgac accagcaaag tgaattatgg tgtaacggta ctgccgacct 840
tcaagggtca accatccaaa ccgttcgttg gcgtgctgag cgcaggtatt aacgcgccca 900
gtccgaacaa agagctggcg aaagagtcc tcgaaaacta tctgctgact gatgaaggtc 960
tggaagcggg taataaagac aaaccctgg gtgccgtagc gctgaagtct tacgaggaag 1020
agttggcgaa agatccacgt attgccgcca ccatgaaaa cgcccagtcc gctttctggt 1080
atgccgtgcg tatcgaagcc cgctgcagg cctcggtcga cgcogaatct agagattata 1140
aagatgacga tgacaaataa taagctagag g 1171

```

```

<210> SEQ ID NO 246
<211> LENGTH: 1171
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Fusion vector

```

```

<400> SEQUENCE: 246
aggaggttct gcatatgaaa ataaaacag gtgcacgcat cctgcatta tccgattaa 60
cgacgatgat gttttccgcc tcggctctcg ccaaaatcga agaaggtaaa ctggtaatct 120
ggattaacgg cgataaaggc tataacggtc tcgctgaagt cggtaagaaa ttcgagaaaag 180
ataccggaat taaagtcacc gttgagcatc cggataaact ggaagagaaa ttcccacagg 240
ttgctgcaac tggcgtatgg cctgacatta tctctgggc acacgaccgc tttggtggct 300
acgctcaatc tggcctggtg gctgaaatca ccccgacaa agcgttcocag gacaagctgt 360
atccgtttac ctgggatgcc gtacgttaca acggcaagct gattgcttac ccgatcgtg 420
ttgaagcgtt atcgtgatt tatacaaaag atctgctgcc gaaccgccca aaaacctggg 480
aagagatccc ggcgctggat aaagaactga aagcgaaaag taagagcgcg ctgatgttca 540
acctgcaaga accgtacttc acctggcccg tgattgctgc tgacgggggt tatgcttca 600
agtatgaaaa cggcaagtac gacattaaag acgtgggctt ggataacgct ggcgcgaaaag 660
cgggtctgac cttcctgggt gacctgatta aaaacaaaca catgaatgca gacaccgatt 720
actccatcgc agaagctgcc ttaataaaag gcgaaacagc gatgaccatc aacggcccgt 780
gggcatggtc caacatcgac accagcaaag tgaattatgg tgtaacggta ctgccgacct 840
tcaagggtca accatccaaa ccgttcgttg gcgtgctgag cgcaggtatt aacgcgccca 900
gtccgaacaa agagctggcg aaagagtcc tcgaaaacta tctgctgact gatgaaggtc 960
tggaagcggg taataaagac aaaccctgg gtgccgtagc gctgaagtct tacgaggaag 1020
agttggcgaa agatccacgt attgccgcca ccatgaaaa cgcccagtcc gctttctggt 1080
atgccgtgcg tatcgaagcc cgctgcagg cctcggtcga cgcogaatct agagattata 1140
aagatgacga tgacaaataa taagctagag g 1171

```

```

<210> SEQ ID NO 247
<211> LENGTH: 392
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

```

-continued

&lt;223&gt; OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 247

```

tagcaggagg ccctgcaggc ctcggtcgac gccgaatcta gaagcgataa aattattcac    60
ctgactgacg acagttttga cacggatgta ctcaaagcgg acggggcgat cctcgtcgat    120
ttctgggcag agtgggtcgg tccgtgcaaa atgatcgccc cgattctgga tgaaatcgct    180
gacgaatata agggcaaaact gaccgttgca aaactgaaca tcgatcaaaa ccctggcact    240
gcgccgaaat atggcatccg tggtatcccg actctgctgc tgttcaaaaa cgggtgaagtg    300
gcggcaacca aagtgggtgc actgtctaaa ggtcagttga aagagaacct ggcggattat    360
aaagatgacg atgacaaata ataagctaga gg                                392

```

&lt;210&gt; SEQ ID NO 248

&lt;211&gt; LENGTH: 426

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 248

```

gaattcaggc gctttttaga ctggtcgtaa tgaaattcag gaggttctgc aggcctcggc    60
cgacgccgaa tctagaagcg ataaaattat tcacctgact gacgacagt ttgacacgga    120
tgtactcaaa gcggacgggg cgatcctcgt cgatttctgg gcagagtgg gcggtccgtg    180
caaaatgatc gccccgattc tggatgaaat cgctgacgaa taccagggca aactgaccgt    240
tgcaaaactg aacatcgatc aaaacacctg cactgcgccg aaatatggca tccgtggtat    300
cccgactctg ctgctgttca aaaacgggtg agtggcggca accaaagtgg gtgcactgtc    360
taaaggctag ttgaaagaga acctggcggg ttataaagat gacgatgaca aataataagc    420
tagagg                                426

```

&lt;210&gt; SEQ ID NO 249

&lt;211&gt; LENGTH: 392

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Fusion vector

&lt;400&gt; SEQUENCE: 249

```

aggaggttct gcaggcctcg gtcgacgccc aatctagaag cgataaaatt attcacctga    60
ctgacgacag ttttgacacg gatgtactca aagcggacgg ggcgatcctc gtcgatttct    120
gggcagagtg gtgctgctcc tgcaaaatga tcgccccgat tctggatgaa atcgctgacg    180
aatatcaggg caaactgacc gttgcaaaac tgaacatcga tcaaaaccct ggcactgctc    240
cgaaatatgg catccgtggt atcccgactc tgctgctggt caaaaacggt gaagtggcgg    300
caaccaaagt ggggtgactg tctaaaggtc agttgaaaga gaacctggcg gattataaag    360
atgacgatga caaataataa gctagaggta cc                                392

```

&lt;210&gt; SEQ ID NO 250

&lt;211&gt; LENGTH: 392

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Fusion vector



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<400> SEQUENCE: 250

```
aggaggttct gcaggcctcg gtcgacgccc aatctagaag cgataaaatt attcacctga 60
ctgacgacag ttttgacacg gatgtactca aagcggacgg ggcgatcctc gtcgatttct 120
gggcagagtg gtgcggtccg tgcaaaatga tcgccccgat tctggatgaa atcgctgacg 180
aatatcaggg caaactgacc gttgcaaaac tgaacatcga tcaaaaccct ggcactgcgc 240
cgaaatatgg catccgtggt atcccgactc tgctgctggt caaaaacggt gaagtggcgg 300
caaccaaaag ggggtgactg tctaaaggtc agttgaaaga gaacctggcg gattataaag 360
atgacgatga caaataataa gctagaggta cc 392
```

<210> SEQ ID NO 251

<211> LENGTH: 528

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 251

```
gaattcagcg gctttttaga ctggtcgtaa tgaattcag gaggttctgc atatgaaaat 60
aaaaacaggt gcacgcattc tcgattatc cgcattaacg acgatgatgt tttccgcctc 120
ggctctcgcc aaaatcatcg aagcccgcct gcaggcctcg gtcgacgccc aatctagaag 180
cgataaaatt attcacctga ctgacgacag ttttgacacg gatgtactca aagcggacgg 240
ggcgatcctc gtcgatttct gggcagagtg gtgcggtccg tgcaaaatga tcgccccgat 300
tctggatgaa atcgctgacg aatatcaggg caaactgacc gttgcaaaac tgaacatcga 360
tcaaaaccct ggcactgcgc cgaaatatgg catccgtggt atcccgactc tgctgctggt 420
caaaaacggt gaagtggcgg caaccaaaag ggggtgactg tctaaaggtc agttgaaaga 480
gaacctggcg gattataaag atgacgatga caaataataa gctagagg 528
```

<210> SEQ ID NO 252

<211> LENGTH: 514

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 252

```
ccatacccgt tttttgggc tagcaggagg cctgcatat gaaaataaaa acaggtgcac 60
gcaccctcgc attatccgca ttaacgacga tgatgttttc cgcctcggct ctcgccaaaa 120
tcacgaagc ccgctgcag gcctcggctg acgcgaatc tagaagcgat aaaattatct 180
acctgactga cgacagtttt gacacggatg tactcaaagc ggacggggcg atcctcgtcg 240
atctctgggc agagtggcgc ggtccgtgca aatgatcgc cccgattctg gatgaaatcg 300
ctgacgaata tcagggcaaa ctgaccgttg caaaactgaa catogatcaa aacctggca 360
ctgcgccgaa atatggcacc cgtggtatcc cgactctgct gctgttcaaa aacggtgaag 420
tggcggcaac caaagtgggt gactgtctta aaggtcagtt gaaagagaac ctggcggatt 480
ataaagatga cgatgacaaa taataagcta gagg 514
```

<210> SEQ ID NO 253

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```

<211> LENGTH: 494
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 253

aggaggttct gcatatgaaa ataaaaacag gtgcacgcat cctcgatta tccgattaa    60
cgacgatgat gttttccgcc tcggctctcg ccaaatcat cgaagccgc ctgcaggcct    120
cggctcagcgc cgaatctaga agcgataaaa ttattcacct gactgacgac agttttgaca    180
cggatgtact caaagcggac ggggcgatcc tcgtcgattt ctgggcagag tggtgcggtc    240
cgtgcaaaaa gatcgccccg attctggatg aaatcgctga cgaatatcag ggcaaaactga    300
ccgttgcaaa actgaacatc gatcaaaacc ctggcactgc gccgaaatat ggcacccgtg    360
gtatcccagc tctgctgctg ttcaaaaacg gtgaagtggc ggcaacccaa gtgggtgcac    420
tgtctaaaag tcagttgaaa gagaacctgg cggattataa agatgacgat gacaaataat    480
aagctagagg tacc                                                    494

```

```

<210> SEQ ID NO 254
<211> LENGTH: 494
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 254

aggaggttct gcatatgaaa ataaaaacag gtgcacgcat cctcgatta tccgattaa    60
cgacgatgat gttttccgcc tcggctctcg ccaaatcat cgaagccgc ctgcaggcct    120
cggctcagcgc cgaatctaga agcgataaaa ttattcacct gactgacgac agttttgaca    180
cggatgtact caaagcggac ggggcgatcc tcgtcgattt ctgggcagag tggtgcggtc    240
cgtgcaaaaa gatcgccccg attctggatg aaatcgctga cgaatatcag ggcaaaactga    300
ccgttgcaaa actgaacatc gatcaaaacc ctggcactgc gccgaaatat ggcacccgtg    360
gtatcccagc tctgctgctg ttcaaaaacg gtgaagtggc ggcaacccaa gtgggtgcac    420
tgtctaaaag tcagttgaaa gagaacctgg cggattataa agatgacgat gacaaataat    480
aagctagagg tacc                                                    494

```

```

<210> SEQ ID NO 255
<211> LENGTH: 1521
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Gene encoding a fusion protein

<400> SEQUENCE: 255

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&lt;210&gt; SEQ ID NO 256

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&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

&lt;220&gt; FEATURE:

&lt;223&gt; OTHER INFORMATION: Gene encoding a fusion protein

&lt;400&gt; SEQUENCE: 256

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&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Artificial Sequence

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<210> SEQ ID NO 258
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<220> FEATURE:
<223> OTHER INFORMATION: Gene encoding a fusion protein

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1. A method of making a pharmaceutical composition comprising a minicell, wherein said minicell displays a membrane protein, wherein said membrane protein is selected from the group consisting of a eukaryotic membrane protein, an archeabacterial membrane protein and an organellar membrane protein.

2. The method of claim 1, wherein said minicell is selected from the group consisting of a eubacterial minicell, a poroplast, a spheroplast and a protoplast.

3. The method of claim 1, wherein said method further comprises adding an adjuvant to said pharmaceutical formulation.

4. The method of claim 1, wherein said method further comprises desiccating said formulation.

5. The method of claim 4, wherein said method further comprises adding a suspension buffer to said formulation.

6. The method of claim 1, wherein said method further comprises making a chemical modification of said membrane protein.

7. The method of claim 6, wherein said chemical modification is selected from the group consisting of glycosylation, deglycosylation, phosphorylation, dephosphorylation and proteolysis.

8. A method of making a pharmaceutical composition comprising a minicell, wherein said minicell displays a membrane protein that is a fusion protein, said fusion protein comprising (i) a first polypeptide, said first polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain; and (ii) a second polypeptide, wherein said second polypeptide is not derived from a eubacterial protein.

9. The method of claim 8, wherein said method further comprises adding an adjuvant to said pharmaceutical formulation.

10. The method of claim 8, wherein said method further comprises desiccating said pharmaceutical formulation.

11. The method of claim 10, wherein said method further comprises adding a suspension buffer to said pharmaceutical formulation.

12. The method of claim 8, wherein said method further comprises making a chemical modification of said membrane protein.

13. The method of claim 12, wherein said chemical modification is selected from the group consisting of glycosylation, deglycosylation, phosphorylation, dephosphorylation and proteolysis.

14. A method of making a pharmaceutical formulation comprising a minicell, wherein said minicell displays a membrane conjugate, wherein said membrane conjugate comprises a membrane component chemically linked to a conjugated compound.

15. The method of claim 14, wherein said method further comprises adding an adjuvant to said pharmaceutical formulation.

16. The method of claim 14, wherein said membrane component is a polypeptide comprising at least one transmembrane domain or at least one membrane anchoring domain, or a lipid that is part of a membrane.

17. The method of claim 14, wherein said conjugated compound is a polypeptide, and the chemical linkage between said membrane compound and said conjugated compound is not a peptide bond.

18. The method of claim 14, wherein said conjugated compound is a nucleic acid.

19. The method of claim 14, wherein said conjugated compound is an organic compound.

20. The method of claim 7, wherein said organic compound is selected from the group consisting of a narcotic, a toxin, a venom, and a sphingolipid.

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